



STIC Search Report

EIC 2800

STIC Database Tracking Number: 129230

TO: Monica Lewis
Location: JEF-5A30
August 12, 2004
Case Serial No. : 10/013,103

From: Jeff Harrison
Location: STIC-EIC2800
JEF-4B68
Phone: 22511

Email: harrison, jeff

Search Notes

Monica,

Attached are the closest of the edited search results from Chemical Abstracts and full-text EP/WO/PCT databases.

Based on this, if you have questions or comments, or if would like refocused searching, please let me know.

Thanks, *Jeff*
Jeff Harrison
Team Leader, STIC-EIC2800
JEF-4B68, 571-272-2511



STIC Search Results Feedback Form

EIC 2800

Questions about the scope or the results of the search? Contact **the EIC searcher or contact:**

Jeff Harrison, EIC 2800 Team Leader
571-272-2511, JEF 4B68

Voluntary Results Feedback Form

➤ *I am an examiner in Workgroup:* *Example: 2810*

➤ *Relevant prior art found, search results used as follows:*

102 rejection
 103 rejection
 Cited as being of interest.
 Helped examiner better understand the invention.
 Helped examiner better understand the state of the art in their technology.

Types of relevant prior art found:

Foreign Patent(s)
 Non-Patent Literature
(journal articles, conference proceedings, new product announcements etc.)

➤ *Relevant prior art not found:*

Results verified the lack of relevant prior art (helped determine patentability).
 Results were not useful in determining patentability or understanding the invention.

Comments:

Drop off or send completed forms to STIC/EIC2800, CP4-9C18



CAS/STN FILE 'HCAPLUS, WPIX' ENTERED AT 14:33:48 ON 12 AUG 2004

L1 2 S US6352940/PN
 L2 SEL PLU=ON L1 1- PRN : 1 TERM
 L3 3 S L2
 L4 SEL PLU=ON L3 1- IC RN : 12 TERMS
 L5 1142392 S L4
 L6 3 S L5 AND L3
 E PASSIVATION/CT
 L7 3118 S "SURFACE TREATMENT"/CT
 L8 43461 S PASSIVATION/CT OR "PASSIVATION KINETICS"/CT OR PASSIVITY/CT OR PASSIVAT#####
 L9 137 S (OXIDE OR INSULAT##### OR DIELEC#####) (3A) (DIRECTLY OR IMMEDIATELY) (4A) SUBSTRATE
 L10 10929 S (OXIDE OR INSULAT##### OR DIELEC#####) (1A) (INITIAL OR FIRST OR BOTTOM OR LOWER OR
 LOWEST OR BELOW)
 L11 213501 S (OXIDE OR INSULAT##### OR DIELEC#####) (1A) (FILM OR LAYER)
 L12 3827 S DIRECTLY (4A) SUBSTRATE
 E OXIDES/CT

FILE 'REGISTRY' ENTERED AT 15:21:03 ON 12 AUG 2004

L14 24489 S O.SI/MF OR O SI/ELF OR SILICA
 L15 396 S N.O.SI/MF OR N O SI/ELF
 E N4 SI3/MF
 L16 3 S "N4 SI3"/MF
 L17 374 S N.SI/MF OR N SI/ELEF
 L18 821 S L17 OR L16 OR N SI/ELF
 E POLYIMID/PCT
 L19 56064 S (POLYIMIDE/PCT OR "POLYIMIDE FORMED"/PCT)
 OR POLYIMIDE OR ((IMIDE OR IMIDO) AND (POLY OR POLYMER OR
 MONOMER OR COPOLYMER OR HOMOPOLYMER))

FILE 'HCAPLUS' ENTERED AT 15:25:50 ON 12 AUG 2004

L20 198369 S L7 OR TREAT##### (2A) SURFACE OR MODIF##### (2A) SURFACE OR (TREAT##### OR
 EXPOS#####) (4A) GAS OR GAS (4A) SURFACE
 L21 9894 S TRILAYER##### OR (TRI OR TRIPLE) (W) LAYER##### OR DISTINGUISH##### (2A) LAYER OR
 SEPARATE (2A) LAYER
 L22 2593 S L21 AND (STACK##### OR SANDWICH##### OR LAMINA##### OR MULTIL? OR (MULTI OR
 MULTIPLE OR THIRD) (2W) (LAYER OR FILM))

FILE 'REGISTRY' ENTERED AT 15:28:41 ON 12 AUG 2004

L23 245 S OZONE OR O3/MF OR O2/MF OR O/MF
 L24 68 S N2/MF OR N/MF OR NITROGEN/CN
 L25 5564 S AMMONIA
 L26 2707 S ARGON

FILE 'HCAPLUS' ENTERED AT 15:29:43 ON 12 AUG 2004

L27 59394 S (L23 OR L24 OR L25 OR L26) (L) GAS
 L28 8277 S (L23 OR L24 OR L25 OR L26) (L) (VAPOR OR VAPOUR)
 L29 27850 S (L23 OR L24 OR L25 OR L26) (L) SURFACE
 L30 35884 S (L23 OR L24 OR L25 OR L26) (L) (TREAT#####)
 OR MODIF#####
 L31 38851 S (L23 OR L24 OR L25 OR L26) (L) (LAYER OR
 FILM)
 L32 334 S (L27 OR L28) AND L29 AND L30
 L33 295 S (L27 OR L28) AND L29 AND L31
 L34 251 S (L27 OR L28) AND L30 AND L31
 L35 1 S L2
 L36 SEL PLU=ON L35 1- IC : 2 TERMS
 L37 SEL PLU=ON L35 1- RN : 9 TERMS
 E INTEGRATED CIRCUITS/CT
 L38 226887 S "INTEGRATED CIRCUITS"/CT OR ("MONOLITHIC MICROWAVE INTEGRATED CIRCUITS"/CT OR
 "OPTICAL INSTRUMENTS (L) CIRCUITS, INTEGRATED"/CT OR "OPTICAL INSTRUMENTS (L) INTEGRATED CIRCUITS"/CT OR "OPTICAL
 INTEGRATED CIRCUITS"/CT OR "OPTICS (L) INTEGRATED"/CT OR "ELECTRONIC PACKAGES"/CT OR MICROELECTRONICS/CT OR
 "MOLECULAR ELECTRONICS"/CT OR "PRINTED CIRCUITS"/CT) OR "SEMICONDUCTOR DEVICES"/CT OR IC OR ICS OR
 CIRCUIT (L) INTEGRATED OR CSP OR SOI OR ELECTRIC CIRCUIT
 L39 6477 S L8 AND (L38 OR H01L?/IC)
 L40 426 S (L9 OR L10 OR L11) AND L12
 L41 5 S L39 AND L40
 L42 7 S (L39 OR L40) AND L7
 L43 6480 S (L39 OR L40) AND L8
 L44 108 S (L39 OR L40) AND L9
 L45 95 S (L39 OR L40) AND L10
 L46 2584 S (L39 OR L40) AND L11
 L47 432 S (L39 OR L40) AND L12
 L48 0 S (L39 OR L40) AND L13
 L49 587 S L14 AND L15 AND (L16 OR L17 OR L18)
 L50 2 S L49 AND L19

L51 15 S L50 OR (L49 AND (POLYIMID#### OR (IMIDE##
 OR IMIDO##)) (2A) (POLY OR POLYMER OR SOFT OR PHOTO OR PHOTORESIST
 OR PHOTOMASK OR LIGHT OR ?DEVELOP? OR PHOTODEFIN##### OR DEFIN#####))
 L52 0 S L49 AND L12
 L53 11 S L49 AND L20
 L54 1 S L49 AND L21
 L55 0 S L49 AND L22
 L56 136 S L49 AND (L23 OR L24 OR L25 OR L26)
 L57 14 S L56 AND L27
 L58 3 S L56 AND GAS(5A)SURFACE
 L59 36 S L20 AND L21 AND L22
 L60 0 S L59 AND PASSIVAT#####
 L61 10 S ((L20 AND (L21 OR L22)) OR (L21 AND L22)) AND PASSIVAT#####
 L62 2 S ((L20 AND (L21 OR L22)) OR (L21 AND L22)) AND REACTIVITY
 L63 25 S ((L20 AND (L21 OR L22)) OR (L21 AND L22)) AND SUSCEPT#####
 L64 0 S ((L20 AND (L21 OR L22)) OR (L21 AND L22)) AND NONREACT?
 L65 5 S ((L20 AND (L21 OR L22)) OR (L21 AND L22)) AND UNREACT?
 L66 1 S ((L20 AND (L21 OR L22)) OR (L21 AND L22)) AND NON REACT#####
 L67 4210 S (L38 OR L39 OR L40 OR L41 OR L42 OR L43 OR
 L44 OR L45 OR L46 OR L47 OR L48 OR L49 OR L50 OR L51 OR L52 OR
 L53 OR L54 OR L55 OR L56 OR L57 OR L58 OR L59) AND DIRECTLY
 L68 751 S (L38 OR L39 OR L40 OR L41 OR L42 OR L43 OR
 L44 OR L45 OR L46 OR L47 OR L48 OR L49 OR L50 OR L51 OR L52 OR
 L53 OR L54 OR L55 OR L56 OR L57 OR L58 OR L59) AND DIRECTLY(6A)SUBSTRATE
 L69 53 S (L38 OR L39 OR L40 OR L41 OR L42 OR L43 OR
 L44 OR L45 OR L46 OR L47 OR L48 OR L49 OR L50 OR L51 OR L52 OR
 L53 OR L54 OR L55 OR L56 OR L57 OR L58 OR L59) AND IMMEDIATE##(6A)SUBSTRATE
 L70 2140 S (L38 OR L39 OR L40 OR L41 OR L42 OR L43 OR
 L44 OR L45 OR L46 OR L47 OR L48 OR L49 OR L50 OR L51 OR L52 OR
 L53 OR L54 OR L55 OR L56 OR L57 OR L58 OR L59) AND (OXIDE OR
 SIO OR SIO2 OR INSULAT####) (3A) (COAT##### OR DEPOSIT#####) (6A)SUBSTRATE
 L71 961 S (L38 OR L39 OR L40 OR L41 OR L42 OR L43 OR
 L44 OR L45 OR L46 OR L47 OR L48 OR L49 OR L50 OR L51 OR L52 OR
 L53 OR L54 OR L55 OR L56 OR L57 OR L58 OR L59) AND (OXIDE OR
 SIO OR SIO2 OR INSULAT####) (2A) (COAT##### OR DEPOSIT#####) (2A)SUBSTRATE
 L72 40 S L71 AND PASSIVAT#####
 L73 527 S (L7 OR L8 OR L9 OR L10 OR L11 OR L12) AND
 (L14 OR L15 OR L16 OR L17 OR L18 OR L19) AND (L20 OR L21 OR
 L22) AND (L23 OR L24 OR L25)
 L74 251 S L73 AND OXIDE(3A) (COAT##### OR LAYER####
 OR FILM##### OR INSULAT#### OR DIELEC#####)
 L75 36 S L73 AND OXIDE(3A)SUBSTRATE
 L76 34 S L74 AND L75
 L77 3 S (L74 OR L75) AND DIRECTLY
 L78 0 S L73 AND OXIDE(3A)FIRST LAYER
 L79 2 S L73 AND OXIDE(3A)FIRST
 L80 0 S L73 AND OXIDE(3A)BOTTOM
 L81 5 S L73 AND OXIDE(3A)LOWER
 L82 15911 S (L7 OR L8 OR L9 OR L10 OR L11 OR L12 OR
 L13) AND (BOND##(3A) (LAYER#### OR FILM) OR (ADHE##### OR GLUE##### OR GLUING))

FILE 'REGISTRY' ENTERED AT 15:52:25 ON 12 AUG 2004

L83 2707 S ARGON

FILE 'HCAPLUS' ENTERED AT 15:53:45 ON 12 AUG 2004

L84	49	S	L82 AND SECOND AND THIRD
L85	251	S	L82 AND L83
L86	1116175	S	L37
L87	3396	S	L82 AND L86
L88	8041	S	L36
L89	135	S	L82 AND L88
L90	1491	S	L82 AND INTERFAC#####
L91	7450	S	L82 AND SURFACE
L92	1803	S	L18(L) (HARD OR PASSIVAT#####)
L93	50	S	L15(L) (ADHE##### OR GLUE##### OR GLUING OR BOND#####)
L94	53	S	SECOND(W) PASSIVAT#####
L95	2278	S	PASSIVAT#####(W) (FILMS OR LAYERS)
L96	2663	S	COMMON(2A) ELEMENT
L97	227	S	L82 AND (L92 OR L93 OR L94 OR L95 OR L96)
L98	220	S	L97 AND PASSIVAT#####
L99	12	S	L97 AND DIRECTLY
L100	87	S	L97 AND SUBSTRATE
L101	80	S	L97 AND OXIDE
L102	30	S	L100 AND L101

L103 1000 S L9 OR (L41 OR L42) OR (L44 OR L45) OR (L50
 OR L51 OR L52 OR L53 OR L54 OR L55 OR L56 OR L57 OR L58 OR L59
 OR L60 OR L61 OR L62 OR L63 OR L64 OR L65 OR L66) OR L69 OR
 L72 OR (L75 OR L76 OR L77 OR L78 OR L79 OR L80 OR L81) OR L84
 OR L89 OR (L93 OR L94) OR (L99 OR L100 OR L101 OR L102)
 L104 167 S L103 AND DIRECTLY
 L105 69 S L103 AND IMMEDIATE##
 L106 605 S L103 AND SUBSTRATE
 L107 75 S L103 AND (MODIF##### OR TREAT#####) (3A) SURFACE
 L108 211 S (L104 OR L105) AND L106
 L109 4 S L107 AND L108
 L110 658 S L103 AND (L104 OR L105 OR L106 OR L107 OR L108 OR L109)
 L111 239 S L110 AND PASSIVAT#####
 L112 471 S L110 AND (IC OR ICS OR INTEGRATED OR CIRCUIT OR SEMICONDUCT##### OR H01L?/IC)
 L113 192 S L111 AND L112
 L114 65 S (L94 OR L95) AND L113
 L115 3 S (L20 OR L21 OR L22) AND L114
 L116 187 S (L111 OR L112) AND (ADHE##### OR GLU##### OR BOND##### (2A) (LAYER OR FILM))
 L117 6 S L116 AND OXIDE(5A)SUBSTRATE
 L118 329 S (L85 OR (L97 OR L98) OR (L104 OR L105 OR
 L106 OR L107 OR L108 OR L109 OR L110 OR L111 OR L112 OR L113
 OR L114 OR L115 OR L116) OR L103 OR (L32 OR L33 OR L34)) AND PASSIVAT#####/TI, IT, ST
 L119 19 S (L104 OR L105) AND L118
 L120 19 S L119 NOT (L109 OR L117 OR L115)
 L121 9 S L120 AND ((L14 OR L15 OR L16 OR L17 OR L18
 OR L19) OR (L23 OR L24 OR L25 OR L26) OR L83 OR ARGON OR GAS (3A) SURFACE)
 L122 2 S L14 AND L15 AND L18 AND L19
 L123 138 S L14 AND L15 AND L18 AND ((L23 OR L24 OR
 L25 OR L26) OR ARGON)
 L124 9 S L123 AND (DIRECTLY OR PASSIVAT#####)
 L125 4 S L123 AND (GLU##### OR ADHE#####)
 L126 13 S (L124 OR L125)
 L127 24 S (L121 OR L122) OR (L109 OR L117 OR L115)
 L128 12 S L126 NOT L127
 L129 92 S L14 AND (L15 OR L18 OR L19) AND PASSIVAT###
 ##### AND (GAS(4A)SURFACE OR (L23 AND L24) OR (L23 AND L25) OR
 (L23 AND L26) OR (L24 AND L25) OR (L24 AND L26) OR (L25 AND L26))
 L130 36 S L126 OR L127
 L131 89 S L129 NOT L130
 L132 89 S L131 AND PASSIVAT#####
 L133 8041 S L36
 L134 7 S L132 AND L133
 L135 1116175 S L37
 L136 6 S H01L023-58?/IC AND OXIDE AND PASSIVAT#####
 L137 0 S H01L023-58?/IC AND OXIDE AND (TRI OR
 TRIPLE OR 3 OR THREE) (W) LAYER#####
 L138 50 S PASSIVAT##### AND OXIDE AND (TRI OR
 TRIPLE OR 3 OR THREE) (W) LAYER#####
 L139 6 S PASSIVAT##### AND OXIDE AND TRILAYER#####
 L140 29 S (L138 OR L139) AND (IC OR ICS OR CIRCUIT OR INTEGRATED OR H01L?/IC)
 L141 0 S L140 AND (COMMON OR DIRECTLY OR IMMEDIATE##
)
 L142 5 S L140 AND SUBSTRATE(4A)(OXIDE OR INSULAT###
 ## OR DIELEC##### OR SIO OR SIO2 OR DIOXIDE OR SILICA)
 L143 0 S L132 AND (L138 OR L139)
 L144 36 S (L126 OR L127)
 L145 22 S (L134 OR L136 OR L139 OR L142) NOT L144
 L146 2 S L145 AND GAS
 L147 4 S L145 AND SURFACE
 L148 4 S L147 NOT L146
 L149 63 S L19(L)PASSIVAT#####
 L150 3106 S L19(L)(PHOTO OR LIGHT OR DEVELOP? OR PHOTOD? OR PHOTO#####)
 L151 9 S L149 AND L150
 L152 80 S (SECOND OR TWO OR 2)(1W)PASSIVAT##### (1W)
 (LAYER##### OR FILM OR COAT##### OR SUBLAYER#####)
 L153 9 S (L19 OR POLYIMIDE OR IMIDE) AND L152
 L154 9 S L153 NOT L151

12aug04 15:26:09 User259284 Session D2866.2

SYSTEM:OS - DIALOG OneSearch
File 348:EUROPEAN PATENTS 1978-2004/Aug W01
(c) 2004 European Patent Office
File 349:PCT FULLTEXT 1979-2002/UB=20040805,UT=20040729
(c) 2004 WIPO/Univentio

Set	Items	Description
S1	13623	DIRECTLY(4N) SUBSTRATE??
S2	3843	(IMMEDIATE?? OR BARE) (4N) SUBSTRATE??
S3	14885	(OXIDE OR INSULAT????? OR OXIDES OR DIOXIDE?? OR SIO2 OR S- IO OR SILICA) (5N) SUBSTRATE??/TI,AB,CM
S4	488	S3 AND PASSIVAT????????/TI,CM,AB
S5	117	S4 AND (IC OR ICS OR INTEGRATED(3N)CIRCUIT??? OR SOI OR CSP OR CHIPSCALE?? OR CHIP()SCALE?? OR MCM OR MCMS OR MULTICHIP?- ?)/TI,AB,CM
S6	5	S5 AND (TRILAYER? OR (TRI OR THREE OR 3)(1W)(LAYER????? OR FILM? ? OR STACK????? OR SANDWICH?????))/TI,AB,CM
S7	244	(SECOND OR 2 OR TWO) (W) PASSIVAT?????(W) (LAYER?? OR FILM?? OR COAT?????)
S8	47	7AND1
S9	8	7AND2
S10	66	7AND3
S11	13	8AND10
S12	57	S8:S11 AND IC=H01L?
S13	1	8AND9
S14	5	9AND10
S15	8	S9 OR S13 OR S14
S16	8	S15 NOT S6
S17	8	S1:S5 AND S16

INSPEC on DIALOG 8/12/2004

Set	Items	Description
S1	3399	PASSIVAT??????(2N) (FILM?? OR LAYER????? OR COAT?????? OR M- ULTI OR MULTIPLE OR TWO OR SECOND OR SUBLAYER???)
S2	25019	'PASSIVATION' OR 'PROTECTIVE COATINGS' OR 'CORROSION PROTE- CTIVE COATINGS'
S3	25454	S1:S2
S4	605	(DIRECT?? OR IMMEDIATE?? OR BARE) (2W) SUBSTRATE
S5	14	3AND4
S6	2	S5 AND SURFACE??(3N) (OXIDE?? OR SIO OR DIOXIDE?? OR SILICA OR SIO2)
S7	3	S5 AND CI=N
S8	4	S5 AND CI=O
S9	0	S5 AND CI=AR
S10	6	S7:S8 NOT S6
S11	3	S10/1999-2004
S12	3	S10 NOT S11
S13	15313	S1:S2 AND PASSIVAT???????????
S14	2	4AND13
S15	1963	(DIRECT?? OR IMMEDIATE?? OR BARE) (4N) SUBSTRATE
S16	17	13AND15
S17	6	S16 AND CI=O
S18	1	S16 AND CI=N
S19	0	S16 AND CI=AR
S20	6	S17:S18
S21	332	S13 AND SUBSTRATE AND OXIDE
S22	207	S21 AND SURFACE
S23	13	S22 AND ADHE???????
S24	0	S22 AND GLU???????
S25	4	S23/1999-2004
S26	9	S23 NOT S25

129230

SEARCH REQUEST FORM Scientific and Technical Information Center - EIC2800

Rev. 3/15/2004 This is an experimental format -- Please give suggestions or comments to Jeff Harrison, JEF-4B68, 272-2511.

Date 8/6/04 Serial # 101013,103 Priority Application Date _____

Your Name M. Harrison Examiner # _____

AU 2829 Phone 219-1838 Room 5A3D

In what format would you like your results? Paper is the default. PAPER DISK EMAIL

If submitting more than one search, please prioritize in order of need. Need before 8/17

The EIC searcher normally will contact you before beginning a prior art search. If you would like to sit with a searcher for an interactive search, please notify one of the searchers.

03-06-04 11:59 AM

Where have you searched so far on this case?

Circle: USPTO DWPI EPO Abs JPO Abs IBM TDB

Other: _____

What relevant art have you found so far? Please attach pertinent citations or Information Disclosure Statements. _____

What types of references would you like? Please checkmark:

Primary Refs Nonpatent Literature Other _____
Secondary Refs Foreign Patents _____
Teaching Refs _____

What is the topic, such as the novelty, motivation, utility, or other specific facets defining the desired focus of this search? Please include the concepts, synonyms, keywords, acronyms, registry numbers, definitions, structures, strategies, and anything else that helps to describe the topic. Please attach a copy of the abstract and pertinent claims.

Claims 17-29

Problem See pages 2 & 3
Solution: " "

Please look for the materials
that are disclosed

Staff Use Only	Type of Search	Vendors
Searcher: <u>HARRISON</u>	Structure (#) _____	STN _____
Searcher Phone: <u>22511</u>	Bibliographic <u>X</u>	Dialog <u>X</u>
Searcher Location: STIC-EIC2800, JEF-4B68	Litigation _____	Questel/Orbit _____
Date Searcher Picked Up: <u>8-11</u>	Fulltext <u>X</u>	Lexis-Nexis _____
Date Completed: <u>8-12-04</u>	Patent Family _____	WWW/Internet _____
Searcher Prep/Rev Time: <u>90</u>	Other _____	Other _____
Online Time: <u>75</u>		

L146 ANSWER 2 OF 2 HCAPLUS COPYRIGHT ACS on STN

AN 1999:783447 HCAPLUS Full-text

DN 132:17954

ED Entered STN: 10 Dec 1999

TI Fabrication of semiconductor devices by plasma CVD of silicon nitride passivation film

IN Miyanaga, Takashi

PA Sony Corp., Japan

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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PI JP 11340223 A2 19991210 JP 1998-141302 19980522

PRAI JP 1998-141302 19980522

AB The plasma CVD of a Si nitride passivation film over a circuit layer on a semiconductor substrate in the title fabrication employs CVD gases including Si nitride-forming reactants and a halo compound gas for involving simultaneous process by vapor depositing of a Si₃N₄ film with the Si nitride-forming reactants and etching of a portion of the depositing Si₃N₄ film with the halo compound gas. The halo compound may be CF₄, C₂F₆, NF₃, SF₆, CHF₃, or ClF₃. The Si nitride-forming reactants may be SiH₄, NH₃, and/or N₂. The simultaneous process gives depositing the Si₃N₄ passivation film an improved coating step coverage for ensuring semiconductor device reliability.

IT Passivation

(film deposition; fabrication of semiconductor devices by plasma CVD of silicon nitride passivation film)

IT 7783-54-2, Nitrogen fluoride (NF₃)

(fabrication of semiconductor devices by plasma CVD of silicon nitride passivation film)

RN 7783-54-2 HCAPLUS

CN Nitrogen fluoride (NF₃) (8CI, 9CI) (CA INDEX NAME)



IT 7664-41-7, Ammonia, reactions 7727-37-9, Nitrogen, reactions 7803-62-5, Silicon hydride (SiH₄), reactions (fabrication of semiconductor devices by plasma CVD of silicon nitride passivation film)

RN 7664-41-7 HCAPLUS

CN Ammonia (8CI, 9CI) (CA INDEX NAME)

NH₃

RN 7727-37-9 HCAPLUS

CN Nitrogen (8CI, 9CI) (CA INDEX NAME)

$N \equiv N$

RN 7803-62-5 HCAPLUS

CN Silane (8CI, 9CI) (CA INDEX NAME)

SiH₄

IT 12033-89-5P, Silicon nitride (Si₃N₄), properties (passivation film, CVD-etching; fabrication of semiconductor devices by plasma CVD of silicon nitride passivation film)

RN 12033-89-5 HCAPLUS

CN Silicon nitride (Si₃N₄) (8CI, 9CI) (CA INDEX NAME)

L148 ANSWER 3 OF 4 HCAPLUS COPYRIGHT ACS on STN

AN 2000:909273 HCAPLUS Full-text

DN 134:65011

ED Entered STN: 28 Dec 2000

TI Integration of low-K SiOF as interlayer dielectric for Al-gapfill application

IN Huang, Richard J.; Iacoponi, John A.

PA Advanced Micro Devices, Inc., USA

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI US 6166427	A	20001226	US 1998-231649	19980115
PRAI US 1998-231649			19980115	

AB A method for producing a dielec. layer in a semiconductor product includes two steps. The 1st step is forming a fluorinated layer (e.g. SiOF or fluorosilicate glass (FSG)) which includes a material formed in part with F. The 2nd step is forming a fill layer (e.g. SiO₂) above the fluorinated layer. The fill layer is substantially free of materials formed in part with F. A top surface of the fill layer can be planarized. Surface treatments and oxide caps can be applied to the planarized surface to form F barriers if part of the fluorinated layer is exposed to higher layers. Such a method, and a semiconductor device or integrated circuit manufactured allow the dielec. constant of an inter-layer dielec. (ILD) to be lowered while also minimizing the complexity and expense of the manufacturing process.

IT Dielectric films
Integrated circuits
Passivation

L128 ANSWER 7 OF 12 HCAPLUS COPYRIGHT ACS on STN

AN 1999:731769 HCAPLUS Full-text

DN 131:331116

ED Entered STN: 17 Nov 1999

TI Reducing bonding pad loss in integrated circuits using a capping layer when etching bonding pad **passivation** openings

IN Hsiao, Yung-Kuan; Wu, Cheng-Ming; Lee, Yu-Hua

PA Taiwan Semiconductor Manufacturing Company, Ltd., Taiwan

PATENT NO. KIND DATE APPLICATION NO. DATE

PI US 5985765 A 19991116 US 1998-75368 19980511

PRAI US 1998-75368 19980511

AB Bonding pad loss is reduced by using a capping layer when contact openings are etched to the bonding pads, while concurrently etching much deeper fuse openings to the substrate. Bonding pads are used on the top surface of integrated circuit semiconductor chips to provide external elec. connections for I/O and power. Fuses are used in the underlying insulating layers to remove redundant defective circuit elements and thereby repair defective chips. It is desirable (cost effective) to etch the contact openings in the **passivation** layer to the bonding pads near the top surface on the chip and to concurrently etch the much deeper fuse openings in the thick underlying insulating layers over the fuses. However, because of the difference in etch depth of the 2 types of openings, the bonding pads composed of Al/Cu are generally overetched, causing bond-pad reliability problems. This invention uses a novel process in which a capping layer, having a low etch rate, is formed on the bonding pads to prevent overetching while the fuse openings are etched to the desired depth in the thicker insulating layers.

IT **Passivation**

(reducing bonding pad loss in integrated circuits using a capping layer when etching openings for bonding pads in **passivation** layers)

IT 12033-89-5, **Silicon nitride, processes** 132614-63-2, **Silicon nitride oxide (Si(N,O))**

(reducing bonding pad loss in integrated circuits using a capping layer when etching openings for bonding pads in **passivation** layers containing)

RN 12033-89-5 HCAPLUS

CN Silicon nitride (Si₃N₄) (8CI, 9CI) (CA INDEX NAME)

RN 132614-63-2 HCAPLUS

CN Silicon nitride oxide (Si(N,O)) (9CI) (CA INDEX NAME)

Component	Ratio	Component Registry Number
N	0 - 1	17778-88-0
O	0 - 1	17778-80-2
Si	1	7440-21-3

IT 7440-37-1, **Argon, processes**

(reducing bonding pad loss in integrated circuits using a capping layer when etching openings for bonding pads in **passivation** layers using gas mixts. containing)

RN 7440-37-1 HCAPLUS

CN Argon (8CI, 9CI) (CA INDEX NAME)

Ar

IT 7631-86-9, **Silica, processes**

(reducing bonding pad loss using a capping layer when etching openings for bonding pads in integrated circuits containing)

RN 7631-86-9 HCAPLUS

CN Silica (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME)

O—Si—O

L121 ANSWER 8 OF 9 HCAPLUS COPYRIGHT ACS on STN

AN 1998:60766 HCAPLUS Full-text

DN 128:187436

ED Entered STN: 02 Feb 1998

TI Plasma deposition of Si-N and Si-O passivation layers
on three-dimensional sensor devices

AU Schmid, P.; Orfert, M.; Vogt, M.

CS TU Dresden, Institut fur Halbleiter- und Mikrosystemtechnik, D-01062,
Dresden, Germany

SO Surface and Coatings Technology (1998), 98(1-3), 1510-1517

CODEN: SCTEEJ; ISSN: 0257-8972

PB Elsevier Science S.A.

DT Journal

LA English

CC 76-14 (Electric Phenomena)

AB In sensor fabrication particularly high demands are made on the **passivation layers**, since the active sensor area is exposed **directly** to the environment. Typical requirements on **passivation layers** are high elec. resistance, high d. against moisture penetration, good adhesion and low mech. stress. In earlier works planar sensors have been successfully **passivated** by plasma-enhanced chemical vapor-deposited (PECVD) SiO and SiN layers. However, many sensor devices are not available in planar techniques, but as three-dimensional (3D) devices. The object of this work was to develop a PECVD **passivation** technique for such 3D sensor devices. For the exptl. work an electron cyclotron resonance (ECR) plasma reactor was used to deposit **passivation layers** on model substrates. Deposition rate and layer quality were measured at various **substrate** locations, orientations and temps. The layer quality was determined by ellipsometer data, IR spectra, SEM and moisture diffusion expts. As result of these investigations the following tendencies could be established. The deposition rate increases in the z-direction (height) by 30% cm⁻¹. At low deposition pressure (0.66-1.33 Pa) the deposition rate depends strongly on the **substrate** orientation, i.e. it decreases from top to side by .apprx.50% and is even lower on the bottom side. Narrow structures with line widths of 1.3 mm and aspect ratios <1 could be well **passivated**. However, narrow undercuts with aspect ratios »1 could not be **passivated** sufficiently. At higher deposition pressures (20.35 Pa), more homogeneous film deposition in gaps and a significantly better coating of bond wires could be achieved.

IT (plasma deposition of Si-N and Si-O passivation
layers on three-dimensional sensor devices)

RN 7631-86-9 HCAPLUS

CN Silica (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME)

O—Si—O

RN 12033-89-5 HCAPLUS

CN Silicon nitride (Si₃N₄) (8CI, 9CI) (CA INDEX NAME)

IT 7664-41-7, Ammonia, reactions 7727-37-9, Nitrogen,
reactions 7782-44-7, Oxygen, reactions 7803-62-5,
Silane, reactions

(plasma deposition of Si-N and Si-O passivation
layers on three-dimensional sensor devices)

RN 7664-41-7 HCAPLUS

CN Ammonia (8CI, 9CI) (CA INDEX NAME)

NH₃

RN 7727-37-9 HCAPLUS

CN Nitrogen (8CI, 9CI) (CA INDEX NAME)

N≡N

RN 7782-44-7 HCAPLUS

CN Oxygen (8CI, 9CI) (CA INDEX NAME)

O—O

L154 ANSWER 2 OF 9 HCAPLUS COPYRIGHT ACS on STN

AN 2000:219098 HCAPLUS Full-text

DN 132:244981

ED Entered STN: 05 Apr 2000

TI Passivation technology combining improved adhesion in passivation and a
scribe street without passivation in semiconductor device fabrication

IN Dass, M. Lawrence A.; Seshan, Krishna; Gaeta, Isaura

PA Intel Corporation, USA

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
-----	---	-----	-----	-----
PI US 6046101	A	20000404	US 1997-1970	19971231

PRAI US 1997-1970 19971231

AB An integrated circuit passivation layer including a first passivation layer portion of silicon nitride treated with nitrous oxide and a **second passivation layer** portion of **polyimide**. Also, a method of passivating an integrated circuit wafer including depositing a first passivation layer over the top surface of an integrated circuit wafer having a scribe street area between adjacent integrated circuit device portions, depositing a **second passivation layer** over the first passivation layer, and patterning the first passivation layer and the **second passivation layer** to expose the scribe street area.

IT Polyimides, uses

(passivation technol. combining improved adhesion in passivation and a
scribe street without passivation in semiconductor device fabrication)

L128 ANSWER 8 OF 12 HCAPLUS COPYRIGHT ACS on STN

AN 1999:686634 HCAPLUS Full-text

DN 131:294446

ED Entered STN: 28 Oct 1999

TI Forming integrated circuit capacitor structures and reducing parasitic capacitance in them

IN Visokay, Mark R.; Colombo, Luigi; McIntyre, Paul; Summerfelt, Scott R.

PA Texas Instruments Incorporated, USA

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI US 5972722	A	19991026	US 1998-60152	19980414
PRAI US 1997-42982P	P	19970414		

AB A high-k dielec. capacitor structure and fabrication method incorporate an adhesion-promoting etch stop layer to promote adhesion of the bottom electrode to the interlevel dielec. layer and to provide a well controlled, repeatable and uniform recess prior to the dielec. deposition. By using a sacrificial layer, e.g. Si₃N₄, this layer can act as an etch stop during the recess etch to eliminate parasitic capacitance between adjacent capacitor cells and can promote adhesion of the bottom electrode material to the substrate.

IT Vapor deposition process

(chemical; of adhesion-promoting sacrificial etch stop layers in integrated circuit capacitor structures)

IT 7631-86-9, Silica, processes 7631-86-9D, Silica, silicon-excess, processes

12033-89-5, Silicon nitride, processes 132614-63-2, Silicon nitride oxide (Si(N,O))

246227-27-0, Silicon nitride (SiN_{0.5-1.34})

(adhesion-promoting sacrificial etch stop layer in integrated circuit capacitor structures)

RN 7631-86-9 HCAPLUS

CN Silica (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME)

0—Si—0

RN 12033-89-5 HCAPLUS

CN Silicon nitride (Si₃N₄) (8CI, 9CI) (CA INDEX NAME)

RN 132614-63-2 HCAPLUS

CN Silicon nitride oxide (Si(N,O)) (9CI) (CA INDEX NAME)

Component	Ratio	Component Registry Number
N	0 - 1	17778-88-0
O	0 - 1	17778-80-2
Si	1	7440-21-3

RN 246227-27-0 HCAPLUS

CN Silicon nitride (SiN_{0.5-1.34}) (9CI) (CA INDEX NAME)

Component	Ratio	Component Registry Number
N	0.5 - 1.34	17778-88-0
Si	1	7440-21-3

IT 7440-37-1, Argon, processes 7782-44-7, Oxygen, processes
(etching by; of integrated circuit capacitor bottom electrodes)

RN 7440-37-1 HCAPLUS

CN Argon (8CI, 9CI) (CA INDEX NAME)

Ar

RN 7782-44-7 HCAPLUS

CN Oxygen (8CI, 9CI) (CA INDEX NAME)

0—O

26/9/1

DIALOG(R) File 2:INSPEC

(c) Institution of Electrical Engineers. All rts. reserv.

6046657 INSPEC Abstract Number: B9811-2530D-027

Title: Adhesion enhancement at Co(P) diffusion barrier/polyimide interfaces

Author(s): O'Sullivan, E.J.; Schrott, A.G.; Sambucetti, C.J.; Kaja, S.; Semkow, K.W.

Author Affiliation: IBM Thomas J. Watson Res. Center, Yorktown Heights, NY, USA

Conference Title: Proceedings of the Symposium on Interconnect and Contact Metallization p.173-81

Editor(s): Rathore, H.S.; Mathad, G.S.; Plougonven, C.; Schuckert, C.C.

Publisher: Electrochem. Soc, Pennington, NJ, USA

Publication Date: 1998 Country of Publication: USA viii+275 pp.

ISBN: 1 56677 184 6 Material Identity Number: XX98-01017

Conference Title: Proceedings of the Symposium on Interconnect and Contact Metallization

Conference Sponsor: Electrochem. Soc

Conference Date: 31 Aug.-5 Sept. 1997 Conference Location: Paris,

Language: English Document Type: Conference Paper (PA)

Treatment: Practical (P); Experimental (X)

Abstract: We studied the adhesion of polyimide to electrolessly deposited Co(P) films used as passivation of Cu lines in multilayer thin film packages. For best adhesion, it was necessary to grow a thin oxide film (<or=60 Å thick) on the Co(P) surface prior to PI application. Oxide films with the desired properties were produced in a well controlled fashion by immersion in a mildly oxidizing borate solution. The oxide thickness and the valency states of Co were investigated ex-situ by means of X-ray photoelectron spectroscopy (XPS). The oxide thickness and the relative abundance of Co³⁺ varied with the pH of the solution. Peel tests (90 degrees) were correlated with XPS analysis of the uncovered substrate and the back of the peeled PI. The results indicated that the interaction of the adhesion promoter with the surface varied with the oxide chemistry, and that the preferred films were Co³⁺ rich. (10 Refs)

Subfile: B

Descriptors: adhesion; chemical interdiffusion; cobalt compounds; diffusion barriers; integrated circuit interconnections; integrated circuit metallisation; integrated circuit reliability; interface structure; mechanical testing; oxidation; passivation; polymer films; surface chemistry; valency; X-ray photoelectron spectra

20/9/4

DIALOG(R) File 2:INSPEC

(c) Institution of Electrical Engineers. All rts. reserv.

6053589 INSPEC Abstract Number: B9811-2560R-073

Title: Demonstration of enhancement-mode p-channel GaAs MOSFETs with $\text{Ga}_{\text{sub}} 2/\text{O}_{\text{sub}} 3/(\text{Gd}_{\text{sub}} 2/\text{O}_{\text{sub}} 3/)$ passivation

Author(s): Ren, F.; Hong, M.; Hobson, W.S.; Kuo, J.M.; Lothian, J.R.; Mannaerts, J.P.; Kwo, J.; Chu, S.N.G.; Chen, Y.K.; Cho, A.Y.

Author Affiliation: Lucent Technol., Bell Labs., Murray Hill, NJ, USA

Conference Title: Proceedings of the Twenty-Sixth State-of-the-Art Program on Compound Semiconductors (SOTAPCOCS XXVI) p.84-90

Editor(s): Buckley, D.N.; Chu, S.N.G.; Hou, H.Q.; Sah, R.E.; Vilcot, J.P.; Deen, M.J.

Publisher: Electrochem. Soc, Pennington, NJ, USA

Publication Date: 1997 Country of Publication: USA ix+322 pp.

ISBN: 1 56677 128 5 Material Identity Number: XX98-00791

Conference Title: Proceedings of the Twenty-Sixth State-of-the-Art Program on Compound Semiconductors (SOTAPCOCS XXVI)

Conference Sponsor: Electrochem. Soc

Conference Date: 4-9 May 1997 Conference Location: Montreal, Que., Canada

Language: English Document Type: Conference Paper (PA)

Treatment: Experimental (X)

Abstract: We report on the first demonstration of an enhancement-mode p-channel GaAs metal oxide semiconductor field effect transistor (MOSFET) directly on GaAs semi-insulating substrate with a high quality $\text{Ga}_{\text{sub}} 2/\text{O}_{\text{sub}} 3/(\text{Gd}_{\text{sub}} 2/\text{O}_{\text{sub}} 3/)$ material as the gate dielectric and ion-implant technology. $\text{Ga}_{\text{sub}} 2/\text{O}_{\text{sub}} 3/(\text{Gd}_{\text{sub}} 2/\text{O}_{\text{sub}} 3/)$ was electron beam deposited from a high purity single crystal $\text{Ga}_{\text{sub}} 2/\text{Gd}_{\text{sub}} 5/\text{O}_{\text{sub}} 12/$ source. The dielectric constant and breakdown field of $\text{Ga}_{\text{sub}} 2/\text{O}_{\text{sub}} 3/(\text{Gd}_{\text{sub}} 2/\text{O}_{\text{sub}} 3/)$ are 14.2 and 3.6 MV/cm, respectively. The source and drain regions were selectively implanted with Zn for low resistance ohmic contacts. AuBe/Pt/Au and Ti/Pt/Au were then deposited for p-ohmic contact and gate electrode, respectively. The device, with a $40*50 \mu\text{m}^2$ gate geometry, exhibits a threshold of -0.6 V, an extrinsic transconductance of 0.3 mS/mm and an excellent gate breakdown field greater than 3 MV/cm. (8 Refs)

Subfile: B

Descriptors: electric breakdown; electron beam deposition; gadolinium compounds; gallium arsenide; gallium compounds; III-V semiconductors; ion implantation; MOSFET; ohmic contacts; passivation; permittivity

L146 ANSWER 1 OF 2 HCAPLUS COPYRIGHT ACS on STN

AN 2000:416633 HCAPLUS Full-text

DN 133:25416

ED Entered STN: 22 Jun 2000

TI Method of forming **passivation** and insulating layers for semiconductor devices using deuterium-containing reaction **gases**

IN Detar, Mark A.

PA Motorola Inc., USA

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
-----	----	-----	-----	-----
PI US 6077791	A	20000620	US 1998-38466	19980311
PRAI US 1996-771352	A2	19961216		

AB Deuterated compds. were used to form **passivation** (20) and other insulating layers to reduce the H content within those films. Semiconductor source **gases**, nitride source **gases**, and dopant **gases** can be obtained in deuterated form. Process steps for forming and etching are substantially the same as those used to form and etch conventional insulating layer. A sintering step can be performed using deuterated **gas** or omitted altogether.

IT Vapor deposition process

(chemical; method of forming **passivation** and insulating layers for semiconductor devices using deuterium-containing reaction **gases**)

IT Dielectric films
Passivation

IT **Gases**

(sintering; method of forming **passivation** and insulating layers for semiconductor devices using deuterium-containing reaction **gases**)

IT 12033-89-5P, Silicon nitride, processes

(method of forming **passivation** and insulating layers for semiconductor devices using deuterium-containing reaction **gases**)

RN 12033-89-5 HCAPLUS

CN Silicon nitride (Si₃N₄) (8CI, 9CI) (CA INDEX NAME)

IT 1590-87-0D, Disilane, deuterated

(method of forming **passivation** and insulating layers for semiconductor devices using deuterium-containing reaction **gases**)

RN 1590-87-0 HCAPLUS

CN Disilane (6CI, 8CI, 9CI) (CA INDEX NAME)

H₃Si—SiH₃

IT 7727-37-9, Nitrogen, uses 7727-37-9D, Nitrogen, compds.,

uses 7782-44-7, Oxygen, uses 10028-15-6, Ozone, uses

(precursor; in method of forming **passivation** and insulating layers for semiconductor devices using deuterium-containing reaction **gases**)

RN 7727-37-9 HCAPLUS

CN Nitrogen (8CI, 9CI) (CA INDEX NAME)

RN 7782-44-7 HCAPLUS

CN Oxygen (8CI, 9CI) (CA INDEX NAME)

RN 10028-15-6 HCAPLUS

CN Ozone (8CI, 9CI) (CA INDEX NAME)

IT 13550-49-7, Ammonia-d3 13587-49-0, Ammonia-d

13780-28-4, Ammonia-d2

26/9/5

DIALOG(R) File 2:INSPEC

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5258325 INSPEC Abstract Number: B9606-2550E-081

Title: Electrical properties of high-quality stacked CdTe/photo-enhanced native oxide for HgCdTe passivation

Author(s): Yan-Kuin Su; Chung-Te Lin; Hsin-Tien Huang; Shou-Jinn Chang; Tai-Ping Sun; Gin-Shiang Chen; Jiunn-Jye Luo

Author Affiliation: Dept. of Electr. Eng., Nat. Cheng Kung Univ., Tainan, Taiwan

Journal: Japanese Journal of Applied Physics, Part 1 (Regular Papers & Short Notes) Conference Title: Jpn. J. Appl. Phys. 1, Regul. Pap. Short Notes (Japan) vol.35, no.2B p.1165-7

Publisher: Publication Office, Japanese Journal Appl. Phys,

Publication Date: Feb. 1996 Country of Publication: Japan

CODEN: JAPNDE ISSN: 0021-4922

SICI: 0021-4922(199602)35:2BL.1165:EPHQ;1-G

Material Identity Number: C579-96005

Conference Title: 1995 International Conference on Solid State Devices and Materials (SSDM '95)

Conference Date: 21-24 Aug. 1995 Conference Location: Osaka, Japan

Language: English Document Type: Conference Paper (PA); Journal Paper (JP)

Treatment: Practical (P); Experimental (X)

Abstract: A novel surface treatment method for obtaining a high-quality CdTe/HgCdTe interface is proposed. By stacking photoenhanced native oxide and CdTe films, we successfully passivated HgCdTe. This technique is advantageous because photoenhanced native oxides can form an excellent interface and adhere to HgCdTe substrate well. Using this novel technique, we have fabricated metal/CdTe/photoenhanced native oxide /HgCdTe structured metal-insulator-semiconductor (MIS) capacitors. From capacitance-voltage (C-V) measurement, we found that the comparison, conventional metal/CdTe/HgCdTe structured MIS capacitors were also fabricated. We found that capacitors with the photoenhanced native oxide layer have a much lower leakage current. Such a marked leakage current reduction is due to the good interfacial properties between the photoenhanced native oxide and the HgCdTe substrate. (11 Refs)

Subfile: B

Descriptors: cadmium compounds; II-VI semiconductors; interface structure; mercury compounds; MOS capacitors; oxidation; passivation; surface treatment

L154 ANSWER 4 OF 9 HCAPLUS COPYRIGHT ACS on STN

AN 1998:277612 HCAPLUS Full-text

DN 128:328873

ED Entered STN: 14 May 1998

TI Radiation imager with discontinuous dielectric

IN Possin, George Edward; Liu, Jianqiang; Kwasnick, Robert Forrest

PA General Electric Company, USA

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
EP 838860	A2	19980429	EP 1997-308466	19971023
EP 838860	A3	19990414		
US 5777355	A	19980707	US 1996-772446	19961223
JP 10214954	A2	19980811	JP 1997-287500	19971021

PRAI US 1996-772446 19961023

AB Radiation imagers having a plurality of photosensitive elements has a two-tier passivation layer disposed between the top patterned common electrode contact layer and resp. photosensor islands. The top passivation layer is a polymer bridge member disposed between adjacent photodiodes so as to isolate defects such as moisture-induced leakage in any bridge island layer to the two adjacent photodiodes spanned by the bridge island.

IT Polyimides, uses
(radiation imager with discontinuous dielec.)

IT 62929-02-6, Probromide 286
(radiation imager with discontinuous dielec.)

RN 62929-02-6 HCAPLUS

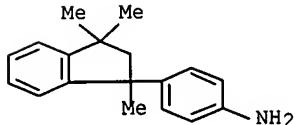
CN 1,3-Isobenzofurandione, 5,5'-carbonylbis-, polymer with 1(or 3)-(4-aminophenyl)-2,3-dihydro-1,3,3(or 1,1,3)-trimethyl-1H-inden-5-amine

CM 1

CRN 60451-10-7

CMF C18 H22 N2

CCI IDS

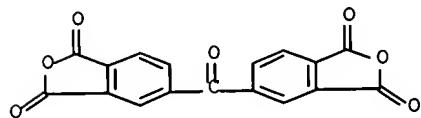


D1-NH2

CM 2

CRN 2421-28-5

CMF C17 H6 O7



L154 ANSWER 5 OF 9 HCAPLUS COPYRIGHT ACS on STN

AN 1998:219317 HCAPLUS Full-text

DN 128:277840

ED Entered STN: 18 Apr 1998

TI Double mask hermetic passivation method providing enhanced resistance to moisture

IN Bryant, Frank R.; Singh, Abha R.; Cunningham, James A.

PA SGS-Thomson Microelectronics, Inc., USA

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI US 5736433	A	19980407	US 1996-778021	19961231
JP 10233454	A2	19980902	JP 1997-353242	19971222

PRAI US 1996-651618 19960522
 US 1996-738738 19961028
 US 1996-778021 19961231

AB A passivation structure is formed using two passivation layers and a protective overcoat layer using two masking steps. The first passivation layer is formed over the wafer and openings are provided to expose portions of the pads for testing the device and fusible links. After testing and laser repair, a second passivation layer is formed over the wafer followed a deposit of the protective overcoat. The protective overcoat is patterned and etched, exposing the pads. The remaining portions of the protective overcoat are used as a mask to remove portions of the second passivation layer overlying the pads. Leads are then attached to pads and the devices are encapsulated for packaging. The second passivation layer overlaps edge portions of the first passivation layer at the bond pads to enhance moisture resistance.

IT Polyimides, processes

(protective overcoat; double mask hermetic passivation method providing enhanced resistance to moisture)

IT 7631-86-9, Silicon dioxide, processes 11105-01-4, Silicon oxynitride

12033-89-5, Silicon nitride, processes

(double mask hermetic passivation method providing enhanced resistance to moisture)

17/TI,PN,PD,AN,AD,IC,AB,AB,K/2 (Item 2 from file: 348)
 DIALOG(R) File 348: (c) European Patent Office. All rts. reserv.

METHOD OF MANUFACTURING A SEMICONDUCTOR THIN FILM TRANSISTOR

PATENT (CC, No, Kind, Date): EP 714140 A1 960529 (Basic)
 EP 714140 A1 980401
 EP 714140 B1 030903
 WO 95034916 951221

APPLICATION (CC, No, Date): EP 95921972 950615; WO 95JP1196 950615

PRIORITY (CC, No, Date): JP 94133374 940615; JP 9572144 950329

INTERNATIONAL PATENT CLASS: H01L-021/205; G02F-001/136; H01L-021/336;
 H01L-021/20; C23C-016/24

ABSTRACT EP 714140 A1

In order to fabricate a high performance thin film semiconductor device using a low temperature process in which it is possible to use low price glass substrates, a thin film semiconductor device has been fabricated by forming a silicon film at less than 450(degree)C, and, after crystallization, keeping the maximum processing temperature at or below 350(degree)C.

In applying the present invention to the fabrication of an active matrix liquid crystal display, it is possible to both easily and reliably fabricate a large, high-quality liquid crystal display. Additionally, in applying the present invention to the fabrication of other electronic circuits as well, it is possible to both easily and reliably fabricate high-quality electronic circuits. (see image in original document)

...SPECIFICATION grown on the substrate in step four. The deposited layer functions as the underlevel protection layer on top of the substrate but functions as a **second passivation layer** on areas in the deposition chamber away from the substrate. Because the underlevel protection layer by itself is able to prevent the diffusion of impurities...SiO₂ film was used as the gate insulator layer and was deposited to a thickness of 1200 Å using PECVD. (Figure 1 (b)) Immediately prior to setting the **substrate** in the PECVD reactor, the substrate was soaked for 20 seconds in a 1.67% dilute hydrofluoric acid solution to remove the native oxide layer...

...SPECIFICATION grown on the substrate in step four. The deposited layer functions as the underlevel protection layer on top of the substrate but functions as a **second passivation layer** on areas in the deposition chamber away from the substrate. Because the underlevel protection layer by itself is able to prevent the diffusion of impurities ...1, a SiO₂) film was used as the gate insulator layer and was deposited to a thickness of 1200 Å using PECVD. (Figure 1 (b)) Immediately prior to setting the **substrate** in the PECVD reactor, the substrate was soaked for 20 seconds in a 1.67% dilute hydrofluoric acid solution to remove the native oxide layer...

...CLAIMS an underlevel protection layer of an insulating material; and a field effect transistor having a semiconductor film formed upon said underlevel protection layer of the **substrate**, a gate insulator layer, and a gate electrode; and an electrically insulating interlevel insulator layer between the interconnects of said field effect transistor;

the formation of a **passivation layer** in said deposition chamber in step 2,

the setting of substrate(s) in said deposition chamber in step 3,

17/TI, PN, PD, AN, AD, IC, AB, AB, K/8 (Item 2 from file: 349)

DIALOG(R) File 349:(c) WIPO/Univentio. All rts. reserv.

CONSTRUCTIONS AND MANUFACTURING PROCESSES FOR THERMALLY ACTIVATED PRINT HEADS

Patent and Priority Information (Country, Number, Date):

Patent: WO 9632267 A1 19961017

Application: WO 96US4855 19960409 (PCT/WO US9604855)

English Abstract

A monolithic printing head having a nozzle configuration in which the heater element is formed using a self-aligned process, where the thickness of the heater, the width of the heater, and the position of the heater in relation to the nozzle are all determined by deposition and etching steps, instead of lithographic processes. In this manner, much greater control of these parameters can be achieved than is generally possible with lithographic processes. No mask is required for the heater. A print head configuration also provides reduced power requirements and incorporates: (1) the provision of a thermally insulating layer between the heater and the substrate; (2) minimizing the thermal mass of the heater and surrounding solid material; (3) minimizing the distance between the heater and the ink meniscus; (4) using a material of relatively high thermal conductivity to **passivate** the heater against corrosion by the ink; and (5) undercutting the substrate in the region of the heater. A method of manufacturing such a nozzle and heater configuration is disclosed.

Detailed Description

... incorporate drive circuitry, data distribution circuitry, and fault tolerance. Also, the active circuitry of the head is protected from chemical attack by the ink by two **passivation layers**: silicon nitride and tantalum

Claim

8. A drop on demand printing head as claimed in claim 7 wherein the layer of material between the heater and the **substrate** is silicon **dioxide**.

9. A drop on demand printing head as claimed in claim 1 further comprising:

- (a) a plurality of drop-emitter nozzles;
- (b) a body of...conductive coating overlying said heater.

19. The invention defined in claim 18 wherein said coating

20. The invention defined in claim 19 further comprising a **passivation** material layer intermediate said heater and said coating.

21. The invention defined in claim 19 wherein said **passivation** layer comprises a tantalum material.

22. A method of fabricating a printing head which includes a self-aligned heater comprising the steps of:

(a) forming...A drop on demand printing head as claimed in claim 31 wherein said actuator is situated on a rim protruding from the surface of said **substrate** in the **immediate** vicinity of said actuator.

33. A drop on demand printing head as claimed in claim 32 characterized in that the substrate material in the region...and the substrate.

37. A drop on demand printing head as claimed in claim 36 wherein the layer of material between the heater and the **substrate** is silicon **dioxide**.

38. A method of manufacture of a drop on demand printing head as claimed in claim 35 wherein the nozzle is formed by anisotropic etching...

L151 ANSWER 1 OF 9 HCAPLUS COPYRIGHT ACS on STN

AN 1996:100848 HCAPLUS Full-text

DN 124:177278

ED Entered STN: 17 Feb 1996

TI Polyimide precursors, photosensitive polymer compositions, and electronic devices prepared therefrom

IN Okabe, Yoshiaki; Ishida, Mina; Miwa, Takao; Takahashi, Akio

PA Hitachi Ltd, Japan; Hitachi Chemical Co Ltd

PATENT NO. KIND DATE APPLICATION NO. DATE

PI JP 07304871 A2 19951121 JP 1994-122020 19940512

PRAI JP 1994-122020 19940512

AB Title precursors comprise repeating units $[A(CO_2H)_2(CONH)(CONHB)]_n$ ($A = 3,3',4,4'$ -biphenyltetrayl, p-terphenyl-3,3",4,4"-tetrayl, 1,2,4,5-benzenetetrayl; B = phenylene or biphenylene; R = C1-3 alkyl; n = 14-270) and photosensitive groups are incorporated to the precursors to give the compns. Thus, 3,3',4,4'-biphenyltracarboxylic dianhydride and tetramethyl-1,4-phenylenediamine were treated in N-methylpyrrolidone to obtain a polyamic acid, 300 g of which was treated with 83.0 g glycidyl methacrylate at 60° for 48 h, and then mixed with 2.6 g 1-phenyl-3-ethoxypropanetrione 2-benzoyloxime and 2.4 g 4,4'-bis(diethylamino)benzophenone, applied on a Si wafer, UV irradiated at 365 nm through a photo mask, developed with choline hydroxide solution, rinsed, and treated at 350° for 30 min to form a polyimide pattern showing T_g 361°.

IT 127669-56-1DP, reaction products with $[(\text{dinitrobenzyl})\text{oxy}]\text{carbonyl}\text{cyclohexylamine}$
174061-93-9P 174061-94-0P 174061-95-1P 174061-96-2P 174061-97-3P

(photoresists containing; polyimide precursors for manufacture of
photosensitive polyimides for passivation films)

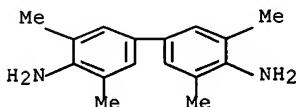
RN 127669-56-1 HCAPLUS

CN [5,5'-Biisobenzofuran]-1,1',3,3'-tetrone, polymer with
3,3',5,5'-tetramethyl[1,1'-biphenyl]-4,4'-diamine (9CI) (CA INDEX NAME)

CM 1

CRN 54827-17-7

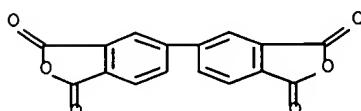
CMF C16 H20 N2



CM 2

CRN 2420-87-3

CMF C16 H6 O6



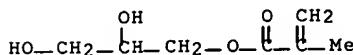
RN 174061-93-9 HCAPLUS

CN [5,5'-Biisobenzofuran]-1,1',3,3'-tetrone, polymer with
2,3,5,6-tetramethyl-1,4-benzenediamine, 2-hydroxy-3-[(2-methyl-1-oxo-2-propenyl)oxy]propyl ester (9CI) (CA INDEX NAME)

CM 1

CRN 5919-74-4

CMF C7 H12 O4



L154 ANSWER 6 OF 9 HCAPLUS COPYRIGHT ACS on STN

AN 1995:820759 HCAPLUS Full-text

DN 123:215940

ED Entered STN: 29 Sep 1995

TI Manufacture apparatus of resin coatings in bevels of semiconductor devices

IN Yamada, Osamu

PA Fuji Electric Co Ltd, Japan

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	JP 07147393	A2	19950606	JP 1994-41653	19940314
	JP 3111794	B2	20001127		
PRAI	JP 1993-178404	A	19930720		
	JP 1993-242263	A	19930929		

AB After etching the bevel regions of a semiconductor substrate to remove the distorted layer, UV radiation using high pressure Hg lamp is applied when the **polyimide** resins (first passivation film) is spread with a contactless resin spreading apparatus, then the silicone resin (second passivation film) is used to mold the final shape. As the adhesion is stronger between the substrate and the first passivation film than between the two passivation films, the first passivation film will survive even there are cracks on the second passivation film due to the pressure on it.

IT Polyimides, processes

Siloxanes and Silicones, processes

(manufacture apparatus of resin coatings in bevels of semiconductor devices)

IT 7440-21-3, Silicon, uses

RL: DEV (Device component use); USES (Uses)

(manufacture apparatus of resin coatings in bevels of semiconductor devices)

L121 ANSWER 9 OF 9 HCAPLUS COPYRIGHT ACS on STN

AN 1995:261571 HCAPLUS Full-text

DN 122:44303

ED Entered STN: 24 Dec 1994

TI Manufacture of **semiconductor** device with aluminum alloy wiring

IN Yamashita, Hiroshi

PA Matsushita Electronics Corp, Japan

SO Jpn. Kokai Tokkyo Koho, 4 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

IC ICM H01L021-3205

CC 76-3 (Electric Phenomena)

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI JP 06267949	A2	19940922	JP 1993-53794	19930315
PRAI JP 1993-53794			19930315	

CLASS

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
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JP 06267949	ICM	H01L021-3205
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AB The title manufacture involves the following steps: (1) forming an interlayer **insulating film** on a **Si substrate** directly or via an elec. **insulating film**, (2) depositing an elec. conductive Al alloy film on the interlayer film by Ar sputtering with heating the **substrate** at 250-300°, (3) etching the elec. conductive film to form a wiring, (4) heating the wiring, (5) depositing a **passivation film** on the wiring, and (6) heating the **passivation film**. Electromigration resistance of Al wiring was improved.

IT 12033-89-5P, Silicon nitride, uses

RL: DEV (Device component use); IMF (Industrial manufacture); PREP (Preparation); USES (Uses)
(**passivation film**, sputtering of; manufacture of **semiconductor** device with aluminum alloy wiring)

IT 7440-21-3, Silicon, uses

RL: DEV (Device component use); USES (Uses)
(**substrate**; manufacture of **semiconductor** device with aluminum alloy wiring)

IT 72893-14-2P

RL: DEV (Device component use); IMF (Industrial manufacture); PEP (Physical, engineering or chemical process); PREP (Preparation); PROC (Process); USES (Uses)
(wiring, sputtering and etching of; manufacture of **semiconductor** device with aluminum alloy wiring)

IT 12033-89-5P, Silicon nitride, uses

RL: DEV (Device component use); IMF (Industrial manufacture); PREP (Preparation); USES (Uses)
(**passivation film**, sputtering of; manufacture of **semiconductor** device with aluminum alloy wiring)

RN 12033-89-5 HCAPLUS

CN Silicon nitride (Si₃N₄) (8CI, 9CI) (CA INDEX NAME)

L154 ANSWER 7 OF 9 HCAPLUS COPYRIGHT ACS on STN

AN 1995:283185 HCAPLUS Full-text

DN 122:44197

ED Entered STN: 10 Jan 1995

TI **Manufacture of compound semiconductor wafers**

IN Yoshimura, Kazunori; Nagayama, Hiroshi

PA Oki Electric Ind Co Ltd, Japan

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI JP 06151583	A2	19940531	JP 1992-296119	19921105
PRAI JP 1992-296119		19921105		

AB In manufacture of compound semiconductor wafers with passivation films covering devices and dicing lines for dividing chips, the dicing line structure comprises passivation films in two sides of the lines and an amorphous film between the two passivation films separated by channels to protect the passivation films from breaking and cracking. The amorphous film can be polyimide resin film or silicon nitride film. In the case of silicon nitride, at least one substrate area along the dicing line has a metal coating.

IT **Polyimides, uses**
(manufacture of compound semiconductor wafers with dicing line structure containing)

IT **12033-89-5, silicon nitride, uses**
(manufacture of compound semiconductor wafers with dicing line structure containing)

L117 ANSWER 5 OF 6 HCAPLUS COPYRIGHT ACS on STN

AN 1994:196003 HCAPLUS Full-text

DN 120:196003

ED Entered STN: 16 Apr 1994

TI Manufacture of photoelectric transfer devices for silicon solar cells

IN Okamoto, Koji; Okuno, Tetsuhiro; Yokozawa, Juji; Moriuchi, Sota; Nakajima, Kazutaka

PA Sharp Kk, Japan

SO Jpn. Kokai Tokkyo Koho, 5 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

IC ICM H01L031-04

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
-----	-----	-----	-----	-----
PI JP 05315628	A2	19931126	JP 1992-116232	19920508
JP 2989373	B2	19991213		
PRAI JP 1992-116232		19920508		

CLASS

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
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JP 05315628	ICM	H01L031-04
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AB The process comprises coating a side (S1) of a Si substrate with a SiO₂-based material, coating another side (S2) with a dopant-containing material, heating the substrate to form a PN bonding layer on S2 and to form an oxide film on S2 and a Si oxide film on S1 simultaneously, and retaining the both films. Passivation films are formed on both sides by the easy process.

IT Photoelectric devices, solar
(silicon, substrates coated with oxide
films for)

IT 7631-86-9, Silica, uses
RL: USES (Uses)
(coatings, on silicon substrates, for solar cells)

IT 7723-14-0, Phosphorus, uses
RL: USES (Uses)
(dopants, oxide coatings containing, on silicon
substrates, for solar cells)

IT 7440-21-3, Silicon, uses
RL: USES (Uses)
(substrates, oxide film formation on, for
solar cells)

17/TI,PN,PD,AN,AD,IC,AB,AB,K/3 (Item 3 from file: 348)
DIALOG(R)File 348:(c) European Patent Office. All rts. reserv.

Conversion of silica precursors to silica at low temperatures.

PATENT (CC, No, Kind, Date): EP 461782 A2 911218 (Basic)
EP 461782 A3 920129
EP 461782 B1 931006

APPLICATION (CC, No, Date): EP 91304882 910530;

PRIORITY (CC, No, Date): US 532828 900604

INTERNATIONAL PATENT CLASS: H01L-021/316; C04B-035/14; C04B-035/64;

ABSTRACT EP 461782 A2

This invention relates to a low temperature method of converting silica precursor coatings to ceramic silica coatings. The method comprises applying a **silica** precursor coating to a **substrate**, exposing the coating to an environment comprising ammonium hydroxide and/or wet ammonia vapors and subjecting the coating to a temperature sufficient to yield the ceramic coating. The methods of the invention are particularly applicable to applying coatings on electronic devices.

...SPECIFICATION infra. Alternatively, the modifying ceramic oxide precursor may be hydrolyzed or partially hydrolyzed, dissolved in the solution comprising the solvent and H-resin and then **immediately** applied to the **substrate**. Various facilitating measures such as stirring or agitation may be used as necessary to produce said solutions.

If compounds of the formula R(sub(x...silicon carbon containing coatings, silicon nitrogen containing coatings, silicon nitrogen carbon containing coatings and/or diamond like carbon coatings.

In a dual layer system, the **second passivation layer** may comprise silicon containing coatings, silicon carbon-containing coatings, silicon nitrogen-containing coatings, silicon carbon nitrogen containing coatings, an additional silicon dioxide and modifying ceramic oxide coating or a diamond-like carbon coating. In a triple layer system, the **second passivation layer** may comprise silicon carbon-containing coatings, silicon nitrogen-containing coatings, silicon carbon nitrogen containing coatings, an additional silicon dioxide and modifying ceramic oxide coating or...

17/TI, PN, PD, AN, AD, IC, AB, AB, K/4 (Item 4 from file: 348)
 DIALOG(R) File 348:(c) European Patent Office. All rts. reserv.

Amine catalysts for the low temperature conversion of silica precursors to silica.

Amin-Katalysatoren fur die Umwandlung von Kieselsaure-Vorprodukten zu Kieselsaure bei niedriger Temperatur.

Catalyseur a base d'amines pour la conversion a temperature basse de produits preliminaires de l'acide silicique en acide silicique.

PATENT (CC, No, Kind, Date): EP 460868 A1 911211 (Basic)
 EP 460868 B1 940518

APPLICATION (CC, No, Date): EP 91304881 910530;

PRIORITY (CC, No, Date): US 532705 900604

INTERNATIONAL PATENT CLASS: C04B-035/14; C04B-041/52;

ABSTRACT EP 460868 A1

This invention relates to a low temperature method of converting coatings of hydrogen silsesquioxane resin or hydrolyzed or partially hydrolyzed R(sub(x))Si(OR)(sub(4-x)) to ceramic silica coatings. The method comprises applying a **silica** precursor coating to a **substrate**, exposing the coating to an environment comprising an **amine** and subjecting the coating to a temperature sufficient to yield the ceramic coating. The methods of the invention are particularly applicable to applying coatings on electronic devices.

...SPECIFICATION Alternatively, the modifying ceramic oxide precursor may be hydrolyzed or partially hydrolyzed, dissolved in the solution comprising the solvent and the H-resin and then **immediately** applied to the **substrate**. Various facilitating measures such as stirring or agitation may be used as necessary to produce said solutions.

If compounds of the formula R(sub(x)...

...sub(4-x)) are to be mixed with modifying ceramic oxide precursors, either or both of these compounds may be hydrolyzed or partially hydrolyzed before or after mixing. For **highly** reactive modifying ceramic oxide precursors such as compounds with propoxide, isopropoxide, butoxide, isobutoxide or acetylacetone substituents, it is preferred that the modifying ceramic oxide precursors...layers, silicon containing coatings, silicon carbon containing coatings, silicon nitrogen containing coatings and/or silicon nitrogen carbon containing coatings.

In a dual layer system, the **second passivation layer** may comprise silicon containing coatings, silicon carbon-containing coatings, silicon nitrogen-containing coatings, silicon carbon nitrogen containing coatings or an additional silicon dioxide and modifying ceramic oxide coating. In a triple layer system, the **second passivation layer** may comprise silicon carbon-containing coatings, silicon nitrogen-containing coatings, silicon carbon nitrogen containing coatings or an additional silicon dioxide and modifying ceramic oxide coating and the third barrier coating may comprise silicon coatings, silicon carbon-containing coatings, silicon nitrogen-containing coatings and silicon carbon nitrogen containing coatings.

The silicon-containing coating described above is applied by a method selected from the group consisting of (a) chemical vapor deposition of a silane, halosilane, halodisilane, halopolysilane or mixtures thereof, (b) plasma enhanced chemical...

L151 ANSWER 4 OF 9 HCPLUS COPYRIGHT ACS on STN

AN 1990:208983 HCPLUS Full-text

DN 112:208983

ED Entered STN: 26 May 1990

TI Surface passivation and barrier height enhancement of n-type indium gallium arsenide (In0.53Ga0.47As) Schottky barrier photodiodes

AU Lee, D. H.; Li, Sheng S.

CS Univ. Florida, Gainesville, FL, 32611, USA

SO Proceedings of SPIE-The International Society for Optical Engineering (1989), 1144(Int. Conf. Indium Phosphide Relat. Mater. Adv. Electron. Opt. Devices, 1st), 174-9

CODEN: PSISDG; ISSN: 0277-786X

DT Journal

LA English

CC 76-5 (Electric Phenomena)

Section cross-reference(s): 73

AB Studies of surface passivation and new barrier height enhancement of n-type In0.53Ga0.47As Schottky barrier photodiodes have been carried out. For surface passivation, various dielec. films such as **SiO₂**, **Si₃N₄** and **polyimide** were studied and compared. The results showed that MSM photodiodes passivated with polyimide film yielded the lowest leakage current, whereas the SiO₂ passivated device had the highest leakage current. A new barrier height enhancement method on n-type In0.53Ga0.47As Schottky diodes was developed by depositing a thin graded superlattice of In0.53Ga0.47As/In0.52Al0.48As (10 periods, 60 Å/per period) on n-In0.53Ga0.47As epilayer using MBE technique. Effective barrier heights of .apprx. 0.71 and .apprx. 0.60 eV were obtained for Au and Cr Schottky contacts deposited on this graded superlattice, resp.

IT 7631-86-9, Silicon dioxide, uses and miscellaneous

12033-89-5, Silicon nitride, uses and miscellaneous

25038-81-7, Dupont PI 2555

(passivation with, of gallium indium arsenide Schottky barrier photodiodes, leakage current in relation to)

IT 25038-81-7, Dupont PI 2555

(passivation with, of gallium indium arsenide Schottky barrier photodiodes, leakage current in relation to)

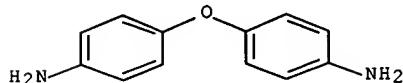
RN 25038-81-7 HCPLUS

CN 1H,3H-Benzo[1,2-c:4,5-c']difuran-1,3,5,7-tetrone, polymer with 4,4'-oxybis[benzenamine] (9CI) (CA INDEX NAME)

CM 1

CRN 101-80-4

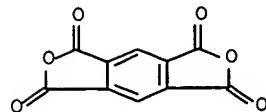
CMF C12 H12 N2 O



CM 2

CRN 89-32-7

CMF C10 H2 O6



L151 ANSWER 3 OF 9 HCAPLUS COPYRIGHT ACS on STN

AN 1991:186959 HCAPLUS Full-text

DN 114:186959

ED Entered STN: 17 May 1991

TI Photosensitive polyimide compositions

IN Okunoyama, Teru; Imagawa, Yasuko

PA Toshiba Chemical Corp., Japan

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI JP 02284924	A2	19901122	JP 1989-108065	19890427
PRAI JP 1989-108065		19890427		

AB The title compns., heat- and chemical-resistant and useful as elec. insulators and passivation films in electronic devices, contain polyimides prepared from 3,3',4,4'-benzophenonetetracarboxylic acid or its dianhydride (I) or alkyl esters and aromatic diamines containing ≥ 90 mol% mixture of $H_2NZ_1(Et)(R_1)Z_2(R_2)(Et)NH_2$ and $CO(Z_1NH_2)_2$ ($Z = CH_2, O, SO_2, CMe_2, C(CF_3)_2, S$; $R_{1-2} = Me, Et, OMe, OEt$; $Z_1 = C_6H_4$, cyclohexylene; ring substituents are all ortho to NH_2). Thus, 232.5 g 4,4'-methylenebis(2,6-diethylaniline), 36.0 g 3,3'-diaminobenzophenone, and 32.2 g I in N-methylpyrrolidone were stirred at 0° for 6 h and then with Ac_2O and pyridine at 100° for 3 h to give 59 g polyimide (II). A cyclohexanone solution of II was coated on a Si wafer, dried, cured by UV through a mask, developed, washed, and dried to give an insulating film with 5% weight loss temperature 495° , glass temperature 295° , and no change after 10 min in C_2HCl_3 , acids, or alkalies; vs. 351, 150, and change, resp., for a conventional photosensitive polyamide.

IT 133396-93-7 133396-94-8 133396-95-9
(photosensitive, chemical- and heat-resistant, for elec. insulation and passivation)

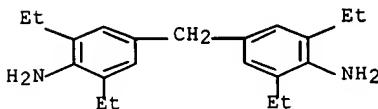
RN 133396-93-7 HCAPLUS

CN 1,3-Isobenzofurandione, 5,5'-carbonylbis-, polymer with
bis(3-aminophenyl)methanone and 4,4'-methylenebis[2,6-diethylbenzenamine]

CM 1

CRN 13680-35-8

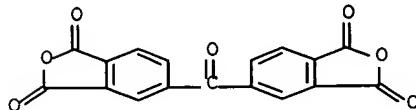
CMF C21 H30 N2



CM 2

CRN 2421-28-5

CMF C17 H6 O7



CM 3

CRN 611-79-0

CMF C13 H12 N2 O

L151 ANSWER 2 OF 9 HCPLUS COPYRIGHT ACS on STN

AN 1991:248595 HCPLUS Full-text

DN 114:248595

ED Entered STN: 28 Jun 1991

TI Photopolymerizable resin compositions for passivation films

IN Kihara, Naoko; Oba, Masayuki; Mikogami, Yukihiro

PA Toshiba Corp., Japan

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI JP 02261862	A2	19901024	JP 1989-83577	19890331
PRAI JP 1989-83577			19890331	

AB Title compns. are prepared polyamide acids prepared from tetracarboxylic dianhydrides and H2NR1NH2 (R1 = divalent organic group containing phenolic OH groups), unsatd. epoxides HCR2:CHR3 (R2 = H, alkyl, aryl; R3 = oxiranyl-terminated organic group), and catalysts. Thus, 21.8 g pyromellitic dianhydride, 21.6 g 4,4'-diaminobiphenyl-3,3'-diol, and N-methylpyrrolidone (I) were stirred at 10° for 5 h and stirred with 12.8 g glycidyl acrylate and 0.7 g Ph3P at 80° for 2 h to give a photocurable composition which was mixed with 3.5 g benzophenone. The mixture was applied on a Si wafer, cured through a neg. photomask in UV light, and soaked in I-MeOH for 5 min to give a good relief pattern.

IT 27082-85-5DP, reaction products with glycidyl acrylate

31669-98-4DP, reaction products with glycidyl acrylate

122962-65-6DP, reaction products with

(glycidoxypropyl) (methacryloyloxypropyl)tetramethyldisiloxane

122988-66-3DP, reaction products with

(glycidoxypropyl) (methacryloyloxypropyl)tetramethyldisiloxane

133830-36-1DP, reaction products with

(glycidoxypropyl) (methacryloyloxypropyl)tetramethyldisiloxane

133830-86-1DP, reaction products with

(glycidoxypropyl) (methacryloyloxypropyl)tetramethyldisiloxane

RL: PREP (Preparation)

(preparation of photocurable, for passivation film on
silicon wafer)

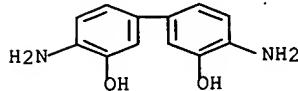
RN 27082-85-5 HCPLUS

CN 1H,3H-Benzo[1,2-c:4,5-c']difuran-1,3,5,7-tetrone, polymer with
4,4'-diamino[1,1'-biphenyl]-3,3'-diol (9CI) (CA INDEX NAME)

CM 1

CRN 2373-98-0

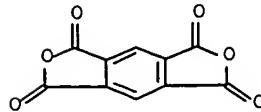
CMF C12 H12 N2 O2



CM 2

CRN 89-32-7

CMF C10 H2 O6



RN 31669-98-4 HCPLUS

L151 ANSWER 5 OF 9 HCPLUS COPYRIGHT ACS on STN

AN 1989:214790 HCPLUS Full-text

DN 110:214790

ED Entered STN: 10 Jun 1989

TI Preparation and properties of soluble and colorless fluorine-containing
polyimide precursors

AU Omote, Toshihiko; Koseki, Kenichi; Yamaoka, Tsuguo

CS Fac. Eng., Chiba Univ., Chiba, 260, Japan

SO *Journal of Photopolymer Science and Technology* (1988), 1(1), 120-1

CODEN: JSTEEW; ISSN: 0914-9244

DT Journal

LA English

CC 42-10 (Coatings, Inks, and Related Products)

Section cross-reference(s): 38, 76

AB In order to form passivation coatings, alpha particle barriers, inter-layer dielecs., etc., by a photo-lithog. technique, four kinds of photosensitive polyimide precursors were prepared by polycondensation of diamines with tetracarboxylic dianhydride followed by esterification with 2-hydroxyethyl methacrylate. The solubility and transparency at 365 nm wavelength were compared to disclose that the presence of CF₃ groups in the amine structure makes the precursor more readily soluble in various organic solvents and optically more transparent. A 3-μm-thick film of the precursor from 2,2-bis(3-amino-4-methylphenyl)hexafluoropropane and biphenyltetracarboxylic dianhydride containing 1 weight% of BP and 5 weight% of MK as photo-initiator provided on a Si wafer gave 0.5 μm L&S patterns with an aspect ratio of 6.0 in spite of neg. working mode.

IT 9043-05-4DP, esters with hydroxyethyl methacrylate

69280-29-1P 120621-33-2P 120720-51-6P

120720-52-7P 120720-53-8P

(preparation of photosensitive, for passivation
coatings, alpha particle barriers and interlayer dielecs.)

RN 9043-05-4 HCPLUS

CN Poly{oxy-1,4-phenyleneimino carbonyl (dicarboxyphenylene) carbonylimino-1,4-phenylene}

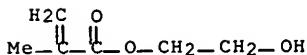
RN 69280-29-1 HCPLUS

CN 1H,3H-Benzo[1,2-c:4,5-c']difuran-1,3,5,7-tetrone, polymer with
4,4'-oxybis[benzenamine], 2-[(2-methyl-1-oxo-2-propenyl)oxy]ethyl ester

CM 1

CRN 868-77-9

CMF C6 H10 O3



CM 2

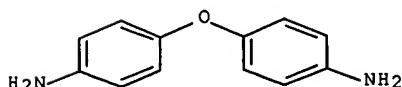
CRN 25038-81-7

CMF (C₁₂ H₁₂ N₂ O . C₁₀ H₂ O₆)_x

CCI PMS

CM 3

CRN 101-80-4

CMF C₁₂ H₁₂ N₂ O

L115 ANSWER 3 OF 3 HCAPLUS COPYRIGHT ACS on STN

AN 1990:563844 HCAPLUS Full-text

DN 113:163844

ED Entered STN: 27 Oct 1990

TI Semiconductor devices coated by passivation
films

IN Miyamoto, Yasunori

PA Matsushita Electric Works, Ltd., Japan

SO Jpn. Kokai Tokkyo Koho, 5 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

IC ICM H01L021-318

ICS H01L021-316

CC 76-3 (Electric Phenomena)

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI JP 02110930	A2	19900424	JP 1988-263777	19881019
PRAI JP 1988-263777			19881019	

CLASS

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
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JP 02110930	ICM	H01L021-318
	ICS	H01L021-316

AB A passivation film coated over a **semiconductor-mounted substrate** comprises Si oxide, Si nitride oxide, and Si nitride layers. The internal stresses of above Si oxide, Si nitride oxide, and Si nitride may be 0.8-1 + 109, 1-2 + 109, and 2-4 + 109 dyn/cm², resp. The 3-layer film prevents formation of cracks during heat-treatment.

IT Semiconductor devices

(passivation films for, triple
layer coating for)

IT 7631-86-9, Silicon oxide (SiO₂), uses and miscellaneous 11105-01-4,
Silicon nitride oxide 12033-89-5, Silicon nitride, properties

RL: USES (Uses)

(passivation film laminated with)

17/TI,PN,PD,AN,AD,IC,AB,AB,K/5 (Item 5 from file: 348)
 DIALOG(R)File 348:(c) European Patent Office. All rts. reserv.

Coatings for microelectronic devices and substrates

Überzugsschicht fur mikroelektronische Anordnungen und Substrate
 Couche de revetement pour dispositifs et substrats micro-electroniques
 PATENT (CC, No, Kind, Date): EP 442632 A2 910821 (Basic)
 EP 442632 A3 920401
 EP 442632 B1 961127

APPLICATION (CC, No, Date): EP 91300701 910130;

PRIORITY (CC, No, Date): US 480399 900215

INTERNATIONAL PATENT CLASS: H01L-021/314; H01L-021/48; H01L-021/312;
 C04B-041/87; C04B-041/89;

ABSTRACT EP 442632 A2

The present invention relates to a method of forming a ceramic or ceramic-like coating on a substrate, especially electronic devices, as well as the substrate coated thereby. The method comprises coating said substrate with a solution comprising a solvent, hydrogen silsesquioxane resin and a modifying ceramic oxide precursor selected from the group consisting of tantalum oxide precursors, niobium oxide precursors, vanadium oxide precursors, phosphorous oxide precursors and boron oxide precursors. The solvent is then evaporated to thereby deposit a preceramic coating on the substrate. The preceramic coating is then ceramified by heating to a temperature of between about 40(degree)C. and about 1000(degree)C. This coating, moreover, may be covered by additional passivation and barrier coatings.

...SPECIFICATION present invention also relates to the formation of additional ceramic or ceramic-like coatings on the coating formed above. In a dual layer system, the **second passivation layer** may comprise silicon containing coatings, silicon carbon-containing coatings, silicon nitrogen-containing coatings, silicon carbon nitrogen containing coatings or an additional silicon dioxide and modifying ceramic oxide coating. In a triple layer system, the **second passivation layer** may comprise silicon carbon-containing coatings, silicon nitrogen-containing coatings, silicon carbon nitrogen containing coatings or an additional silicon dioxide and modifying ceramic oxide coating...Alternatively, the modifying ceramic oxide precursor may be hydrolyzed or partially hydrolyzed, dissolved in the solution comprising the solvent and the H-resin and then immediately applied to the **substrate**. Various facilitating measures such as stirring or agitation may be utilized as necessary to produce said solutions.

The preceramic solution may optionally be catalyzed by...layers, silicon containing coatings, silicon carbon containing coatings, silicon nitrogen containing coatings and/or silicon nitrogen carbon containing coatings.

In a dual layer system, the **second passivation layer** may comprise silicon containing coatings, silicon carbon-containing coatings, silicon nitrogen-containing coatings, silicon carbon nitrogen containing coatings or an additional silicon dioxide and modifying ceramic oxide coating. In a triple layer system, the **second passivation layer** may comprise silicon carbon-containing coatings, silicon nitrogen-containing coatings, silicon carbon nitrogen containing coatings or an additional silicon dioxide and modifying ceramic oxide coating...

...SPECIFICATION present invention also relates to the formation of additional ceramic or ceramic-like coatings on the coating formed above. In a dual layer system, the **second passivation layer** may comprise silicon-containing coatings, silicon-carbon-containing coatings, silicon-nitrogen-containing coatings, silicon-carbon-nitrogen-containing coatings or an additional silicon dioxide and modifying ceramic oxide coating. In a triple layer system, the **second passivation layer** may comprise

silicon-carbon-containing coatings, silicon-nitrogen-containing coatings, silicon-carbon-nitrogen-containing coatings or an additional silicon dioxide and modifying ceramic oxide coating...Alternatively, the modifying ceramic oxide precursor may be hydrolysed or partially hydrolysed, dissolved in the solution comprising the solvent and the H-resin, and then **immediately** applied to the **substrate**.

Various facilitating measures such as stirring or agitation may be utilized as necessary to produce said solutions.

The preceramic solution may optionally be catalysed by...layers, silicon-containing coatings, silicon-carbon-containing coatings, silicon-nitrogen-containing coatings and/or silicon-nitrogen-carbon-containing coatings.

In a dual layer system, the **second passivation layer** may comprise silicon-containing coatings, silicon-carbon-containing coatings, silicon-nitrogen-containing coatings, silicon-carbon-nitrogen-containing coatings or an additional silicon dioxide and modifying ceramic oxide coating. In a triple layer system, the **second passivation layer** may comprise silicon-carbon-containing coatings, silicon-nitrogen-containing coatings, silicon-carbon-nitrogen-containing coatings or an additional silicon dioxide and modifying ceramic oxide coating...

17/TI,PN,PD,AN,AD,IC,AB,AB,K/6 (Item 6 from file: 348)
 DIALOG(R) File 348:(c) European Patent Office. All rts. reserv.

MOS field-effect transistor and method of making the same.

PATENT (CC, No, Kind, Date): EP 255133 A2 880203 (Basic)
 EP 255133 A3 881207
 EP 255133 B1 931006

APPLICATION (CC, No, Date): EP 87111043 870730;

PRIORITY (CC, No, Date): JP 86178889 860731

INTERNATIONAL PATENT CLASS: H01L-029/10; H01L-029/784;

ABSTRACT EP 255133 A2

The present invention relates to a semiconductor device comprising a semiconductor substrate (1) of a first conductivity type or an insulator, a source (4) comprising an impurity layer of a second conductivity type disposed on said semiconductor **substrate** or said **insulator**, a drain (5) comprising an impurity layer of the second conductivity type disposed on said semiconductor **substrate** or said **insulator**, an impurity layer (6) of the first conductivity type formed between said source and said drain, a gate (3) formed on said impurity layer of the first conductivity type via an insulation film, and an impurity layer (7) of the second conductivity type having an impurity concentration lower than that of said source and said drain, said impurity layer of the second conductivity type being disposed between said source, said drain and said impurity layer of the first conductivity type, and said semiconductor substrate of the first conductivity type or said insulator.

...SPECIFICATION layer 2 located under the gate 3 by the gate voltage $V_{sub(GS)}$. Accordingly, a channel is formed on the surface of the semiconductor **substrate** 1 **immediately under the insulation layer** 2. When the illustrated MOSFET is turned ON, therefore, the drain current is distributed so as to be concentrated to a range of several nm (ten angstroms) in depth from the...thickness of 800 nm (8,000 Å) and worked to have a wiring shape by the photolithography technique. The electrode 9 is formed. Lastly the **passivation film** 11 is deposited.

By means of the processing steps heretofore described, the MOSFET according to the present invention explained by referring to Figs. 2 to

...

...CLAIMS a first conductivity type (p) or an insulator;
 a source (4) comprising an impurity layer of a second conductivity type (n) disposed on said semiconductor **substrate** or said **insulator**;
 a drain (5) comprising an impurity layer of the second conductivity type disposed on said semiconductor **substrate** or said **insulator**;
 an impurity layer (6) of the first conductivity type formed between said source and said drain;
 a gate (3) formed on said impurity layer of...

...Claim 1, wherein said semiconductor substrate of the first conductivity type has such a structure that a monocrystalline thin semiconductor film is formed on an **insulator substrate**.

3. A fabrication method of semiconductor device comprising the steps of:

(1) forming an oxide film (2) on the surface of a semiconductor substrate (1...)

...the surface of an insulator;

(2) forming an impurity layer (7) by implanting impurity ions of a second conductivity type from the top of said **oxide** film into said **substrate** or said **insulator**;
 (3) forming a polysilicon gate (3) on said oxide film;
 (4) forming a source (4) and a drain (5) by implanting impurity ions of the...

L154 ANSWER 8 OF 9 HCAPLUS COPYRIGHT ACS on STN

AN 1988:503163 HCAPLUS Full-text

DN 109:103163

ED Entered STN: 17 Sep 1988

TI Passivation of semiconductor devices with polyimides

IN Eguchi, Masuichi; Hiramoto, Yoshi; Manabe, Shinichi

PA Toray Industries, Inc., Japan

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI JP 63015424	A2	19880122	JP 1986-160172	19860708
JP 07116407	B4	19951213		
PRAI JP 1986-160172		19860708		

AB A polyimide-type varnish, consisting of polyamic acids having a structural repeating unit $\text{COR}_1(\text{CO}_2\text{H})_m\text{CONHR}_2\text{NH}$ (R_1 = tri- or tetravalent organic group; R_2 = divalent organic group; $m = 1, 2$), tertiary amines, and $\text{R}_3\text{nSi}(\text{OH})_4-n$ (R_3 = monovalent organic group; $n = 1, 2, 3$) or their partial condensates, is applied on exposed parts of a p-n junction and then heat-treated for passivation of semiconductor devices. Semiconductor devices having high reliability can be prepared Benzophenonetetracarboxylic acid dianhydride-bis(3-aminopropyl)tetramethyldisiloxane-diaminodiphenyl ether-pyromellitic acid dianhydride copolymer was mixed with dimethylaminoethyl methacrylate and OCD Type 2 (silanol compound), applied on a p-n junction of a power diode, and heat-treated to give a passivation coating having excellent resistance to exposure in pressurized saturated vapor at 120°.

IT 116164-52-4
(passivation coating containing, for semiconductors)

RN 116164-52-4 HCAPLUS

CN 1H, 3H-Benzo[1,2-c:4,5-c']difuran-1,3,5,7-tetrone, polymer with 5,5'-carbonylbis[1,3-isobenzofurandione], ar,ar'-oxybis[benzenamine] and 3,3'-(1,1,3,3-tetramethyl-1,3-disiloxanediyl)bis[1-propanamine]

CM 1

CRN 27133-88-6

CMF C12 H12 N2 O

CCI IDS



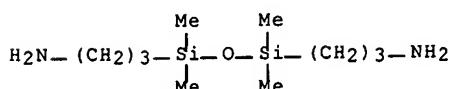
D1-NH2

1/2 (D1-O-D1)

CM 2

CRN 2469-55-8

CMF C10 H28 N2 O Si2



CM 3

6/TI, PN, PD, AN, AD, IC, AB, AB, K/2 (Item 2 from file: 348)
 DIALOG(R) File 348: (c) European Patent Office. All rts. reserv.

A trench-incorporated monolithic semiconductor capacitor and high density dynamic memory cells including the capacitor.

PATENT (CC, No, Kind, Date): EP 220392 A2 870506 (Basic)
 EP 220392 A3 890614
 EP 220392 B1 930505

APPLICATION (CC, No, Date): EP 86110459 860729;

PRIORITY (CC, No, Date): US 792996 851030

INTERNATIONAL PATENT CLASS: H01L-027/10; H01L-021/82;

ABSTRACT EP 220392 A2

A high density **integrated circuit** structure, for example a dynamic memory cell, is described which includes an active/passive device in combination with a capacitor structure. The capacitor structure is of the polysilicon-oxide-silicon type and is formed on the sidewalls of a mesa-shaped and dielectrically isolated region of silicon material resulting from the formation of an isolation trench in the silicon. The trench is filled with a plastic material, such as polyimide. The capacitor is formed by the isolated region of silicon material (14) which functions as the first capacitor plate, a doped polysilicon layer (22a) provided on the vertical walls of the mesa serving as the second capacitor plate and a thin dielectric layer (21) interposed between the two plates serving as the capacitor's dielectric. Since the polysilicon is wrapped around the periphery of the mesa as a coating on the vertical sidewalls thereof, it gives rise to a large storage capacitance without an increase in the cell size.

...CLAIMS serving as the dielectric for said capacitor structure;

d) forming a thin layer (22) of doped polysilicon on the resulting structure;
 e) forming a **dual passivating layer** of silicon dioxide and silicon nitride (23, 24), in order, on the resulting structure;
 f) removing by using a photolithographic process the nitride-oxide...

...j) making ohmic contacts (25a, 29a) with both said polysilicon tab and the reach-through region.

2. The method of claim 1 further including the **step** of filling said trenches for device isolation purposes with a material (19a) selected from a group consisting of polyimide, polysilicon and silicon dioxide.
3. The method of claim 2 wherein said substrate is of the P- type of conductivity.
4. The method of claim 3 wherein said epitaxial layer...

...capacitor.

7. The capacitor structure of claim 6 wherein said insulating layer is silicon dioxide.

8. The capacitor structure of claim 6 further comprising a **passivation** layer covering said conductive polysilicon.

9. The capacitor structure of claim 8 wherein said **passivation** layer is comprised of a thin silicon dioxide layer formed by thermal reoxidation of the polysilicon layer and a silicon nitride overcoat layer.

10. The...location where a capacitor is to be formed;

b) a pattern of substantially vertical isolation trenches extending from one surface of said structure into said **substrate** through said epitaxial layer, thereby delineating a plurality of mesa shaped isolated regions of semiconductor material;

c) a conductive polysilicon layer provided at least partially on the vertical sidewalls of said isolated regions where a capacitor structure is needed thereby forming the first...

...of the second electrode plate of the capacitor.

18. A high density memory according to claim 17 wherein said active device is either a vertical **NPN** bipolar transistor or **P-FET**.

19. A high density memory according to claim 13 wherein said substrate is of the **P⁺** type of conductivity and said epitaxial layer is

...
...density memory according to claim 13 wherein said insulating layer is silicon dioxide.

24. A high density memory according to claim 23 further comprising a **passivation** layer formed on said polysilicon layer.

25. A high density memory according to claim 24 wherein said **passivation** layer is comprised of a thin silicon dioxide layer formed by thermal reoxidation of the polysilicon layer and a silicon nitride overcoat layer.

26. A...

...is material selected from the group consisting of polyimide, polysilicon and silicon dioxide.

28. A high density memory according to claim 13 wherein the configuration of the base of said mesa-shaped isolated region is selected from the group of configurations consisting of rectangular, square, circular, serpentine, and comb-like configuration...

...including a conductive polycrystalline silicon layer on the sidewall of said trench and said regions which said polycrystalline silicon layer is separated from a semiconductor **substrate** by the capacitor silicon **dioxide** dielectric layer;

an extended portion of said polycrystalline silicon layer extends to the top surface of said regions to act as an contact to said...

layer serving as the dielectric for said capacitor structure;

d) forming a thin layer of doped polysilicon on the resulting structure;

e) forming a dual **passivating** layer of silicon dioxide and silicon nitride, in order, on the resulting structure;

f) removing by using a photolithographic process the nitride-oxide-polysilicon composite...

...CLAIMS structure de condensateur,

d) la formation d'une couche mince (22) de polysilicium dope sur la structure obtenue,

e) la formation d'une couche de **passivation** double de dioxyde de silicium et de nitrure de silicium (23, 24), afin d'effectuer les opérations suivantes sur la structure obtenue,

f) le retrait...

L151 ANSWER 9 OF 9 HCPLUS COPYRIGHT ACS on STN

AN 1983:5047 HCPLUS Full-text

DN 98:5047

ED Entered STN: 12 May 1984

TI Siloxanes

IN Berger, Abe

PA M and T Chemicals Inc. , USA

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI EP 54426	A2	19820623	EP 1981-305864	19811214
EP 54426	A3	19820811		
US 4395527	A	19830726	US 1980-216599	19801215

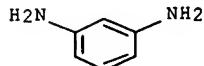
AB Monomers or polymers containing the linkage $ZZ1Z2SiR2(OSiR12)x(OSiR2R3)y(OSiR4 R5)xOSiR2Z2Z1Z$ (Z = substituted or unsubstituted aromatic compound; $Z1$ = O, S, SO, SO₂, SO₂NH, NH₂SO₂, CONH, NHCO, CO₂, O₂C; $Z2$ = substituted or unsubstituted hydrocarbylene; R, R₁, R₂, R₃, R₄, R₅ = substituted or unsubstituted hydrocarbyl; x, y, z = 0-100) were prepared and used in modification of polyimides and other polymers. Thus, a mixture consisting of 50% aqueous NaOH 43.28, DMSO 112, PhMe 120, and p-aminophenyl [123-30-8] 59.95 parts was heated under N, azeotropically distilled to removed water, stirred 7-8 h while the temperature increased to 122°, cooled to .apprx.80°, treated with 86.6 parts bis(chlorobutyl)tetramethyldisiloxane [72066-91-2] dropwise to maintain the reaction temperature at .apprx.80°, heated .apprx.16 h at 80°, and distilled at 295-300° at 0.5-2 mm Hg to give bis(p-aminophenoxybutyl)tetramethyl disiloxane (I) [72066-92-3] which was a colorless liquid which eventually solidified to a white solid with melting 48-49°. A mixture consisting of I 54.64, m-phenylenediamine 29.94, and n-methylpyrrolidone 636 g was cooled to 0°, treated portionwise over a 4-h period with 127.05 g benzophenone tetracarboxylic dianhydride, stirred 10 h at room temperature to give a dark amber clear viscous solution of the corresponding poly(half-amide) which was coated on a glass slide to .apprx.0.2 mil thickness, heated 2 h at 120°, heated 2 h at 135°, heated 2 h at 185°, heated 2 h at 250°, and heated 0.5 h at 300° to give a polyimide [83874-52-6] coating which bonded tenaciously to the glass slide even after immersion in boiling water for 6 h. The polyimide was excellent as a **passivation and/or protective coating for semiconductor devices** including application of the material to exposed portions of P-N junctions. Low leakage current <0.3 μamps was observed at 0.31 μamps and 2000 V. The polyimide could resist 450° for <1 h.

IT 83874-61-7D, reaction products with azidodinnamyl chloride
(photosensitive)

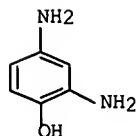
RN 83874-61-7 HCPLUS

CN 1,3-Isobenzofurandione, 5,5'-(1-methylethylidene)bis(4,1-phenyleneoxy)bis-, polymer with 1,3-benzenediamine and 2,4-diaminophenol

CM 2

CRN 108-45-2
CMF C6 H8 N2

CM 3

CRN 95-86-3
CMF C6 H8 N2 O

L151 ANSWER 6 OF 9 HCAPLUS COPYRIGHT ACS on STN

AN 1985:185995 HCAPLUS Full-text

DN 102:185995

ED Entered STN: 02 Jun 1985

TI Organic solvent-soluble photosensitive polyimides

PA Ube Industries, Ltd., Japan; Nippon Telegraph and Telephone Public Corp.

SO Jpn. Kokai Tokkyo Koho, 6 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

IC ICM C08G073-12

CC 37-3 (Plastics Manufacture and Processing)

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI JP 59232122	A2	19841226	JP 1983-106561	19830614
PRAI JP 1983-106561			19830614	

CLASS

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
JP 59232122	ICM	C08G073-12

AB Polyimides useful as elec. insulating materials and passivation membranes are copolycondensates of a biphenyltetracarboxylic acid or its dianhydride and aromatic diamines H₂NC₆H₄CH:CHCOR (R = C₆H₄NH₂, CH:CHC₆H₄NH₂) and (H₂NC₆H₄)₂Z (Z = O, CO, p-OC₆H₄SO₂C₆H₄O-p). Thus, 2,3,3',4'-biphenyltetracarboxylic dianhydride 1580, 3,3'-diaminodibenzalacetone 710, and 4,4'-diaminodiphenyl ether 538 mg were mixed in N-methyl-2-pyrrolidone (I) to give a polyamic acid which was dissolved in I and imidated using Ac₂O and pyridine to give a polyimide [96250-25-8] having inherent viscosity (30°) 0.45, good film formability, solubility in I 10.0%, and heat decomposition temperature 500°. The polyimide film (1.0 μ) was cured by light irradiation of 1.3 J/cm².

IT 96250-23-6P 96250-24-7P 96250-25-8P

(preparation and photochem. crosslinking of)

IT 96250-23-6P 96250-24-7P 96250-25-8P

(preparation and photochem. crosslinking of)

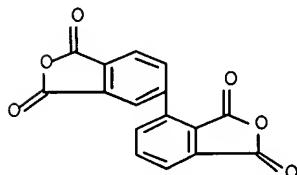
RN 96250-23-6 HCAPLUS

CN [4,5'-Biisobenzofuran]-1,1',3,3'-tetrone, polymer with 1-(4-aminophenyl)-3-(3-aminophenyl)-2-propen-1-one and 4,4'-oxybis[benzenamine] (9CI) (CA INDEX NAME)

CM 1

CRN 36978-41-3

CMF C16 H6 O6



CM 2

CRN 30278-77-4

CMF C15 H14 N2 O

L151 ANSWER 8 OF 9 HCAPLUS COPYRIGHT ACS on STN

AN 1983:90599 HCAPLUS Full-text

DN 98:90599

ED Entered STN: 12 May 1984

TI Characterization of a polyimide passivation on thin film resistor networks

AU Scheiman, Gerald R.

CS Natl. Semiconduct., Santa Clara, CA, USA

SO International Journal for Hybrid Microelectronics (1982), 5(2), 202-4

CODEN: IMICDJ; ISSN: 0277-8270

DT Journal

LA English

CC 38-2 (Plastics Fabrication and Uses)

Section cross-reference(s): 76

AB Under certain conditions, Hitachi PIQ [55478-71-2] is an excellent passivation material for thin film resistor networks. Adhesion, chemical stability, and wire bond shorting protection are comparable or superior to those of SiO₂. PIQ's lithog. and etch properties are within the tolerance required for most thin film circuits. The only drawback to its use is in laser trimmings. PIQ absorbs significant amts. of energy from the 106.4 nm light used in most laser trim systems, which precludes its use for most laser-trimmed resistors.

IT 55478-71-2

(passivation by, on elec. resistors)

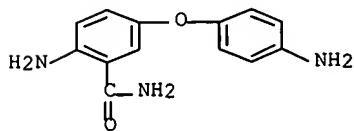
RN 55478-71-2 HCAPLUS

CN Benzamide, 2-amino-5-(4-aminophenoxy)-, polymer with 1H,3H-benzo[1,2-c:4,5-c']difuran-1,3,5,7-tetrone, 5,5'-carbonylbis[1,3-isobenzofurandione] and 4,4'-oxybis[benzenamine] (9CI) (CA INDEX NAME)

CM 1

CRN 40763-98-2

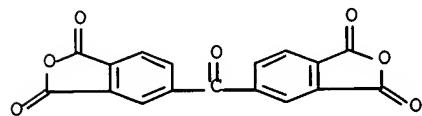
CMF C13 H13 N3 O2



CM 2

CRN 2421-28-5

CMF C17 H6 O7



CM 3

CRN 101-80-4

CMF C12 H12 N2 O

L151 ANSWER 7 OF 9 HCAPLUS COPYRIGHT ACS on STN

AN 1983:523743 HCAPLUS Full-text

DN 99:123743

ED Entered STN: 12 May 1984

TI Selective etching of polyimide type resin film

IN Saiki, Atsushi; Iwayanagi, Takao; Nonogaki, Saburo; Nishida, Takashi;
Harada, Seiki

PA Hitachi, Ltd. , Japan

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI EP 82417	A2	19830629	EP 1982-111392	19821209
EP 82417	A3	19860326		
EP 82417	B1	19890315		
JP 58108229	A2	19830628	JP 1981-205326	19811221
JP 02019972	B4	19900507		
US 4436583	A	19840313	US 1982-445576	19821130

PRAI JP 1981-205326 19811221

CLASS

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
EP 82417	IC	H01L021-312

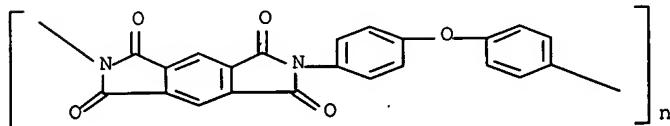
AB A selective etching method for polyimide resin films useful as interlevel insulation for multilevel metalization or as passivation films of transistors, integrated circuits, and large-scale integrated circuits comprises using an etching mask consisting of a neg.-type photoresist material prepared by adding a photosensitive reagent to an unsatd. ketone polymer as the base resin and an etching solution containing 20-40% hydrazine hydrate and 60-80% of a polyamine. Thus, 2,6-bis(4-azidobenzylidene)-4-methylcyclohexanone [5284-79-7] photosensitive reagent was added to poly(isopropenyl Me ketone) [25988-32-3] to give a neg.-type photoresist which was dissolved in cyclohexanone, coated on a 2- μ thick PIQ [55478-71-2] film, and prebaked 20 min at 85°. A photomask having aperture pattern \geq 2 μ square was placed in intimate contact with the film and irradiated with UV light. The film was developed with cyclohexanone, rinsed with Bu acetate, and baked 20 min at 140°. The developed film was etched 20 min at 30° with a 3:7 mixture of hydrazine hydrate and ethylenediamine [107-15-3].

IT 25036-53-7 55478-71-2

(films, etching of, selective, with photoresists)

RN 25036-53-7 HCAPLUS

CN Poly[(5,7-dihydro-1,3,5,7-tetraoxobenzo[1,2-c:4,5-c']dipyrrole-2,6(1H,3H)-diyl)-1,4-phenyleneoxy-1,4-phenylene] (9CI) (CA INDEX NAME)



RN 55478-71-2 HCAPLUS

CN Benzamide, 2-amino-5-(4-aminophenoxy)-, polymer with 1H,3H-benzo[1,2-c:4,5-c']difuran-1,3,5,7-tetrone, 5,5'-carbonylbis[1,3-isobenzofurandione] and 4,4'-oxybis[benzenamine] (9CI) (CA INDEX NAME)

CM 1

CRN 40763-98-2

CMF C13 H13 N3 O2

L154 ANSWER 9 OF 9 HCPLUS COPYRIGHT ACS on STN

AN 1983:171768 HCPLUS Full-text

DN 98:171768

ED Entered STN: 12 May 1984

TI Magnetic-bubble memory device

PA Nippon Electric Co., Ltd., Japan

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
-----	-----	-----	-----	-----

PI JP 57167191 A2 19821014 JP 1981-50092 19810403

PRAI JP 1981-50092 19810403

AB The fabrication of a magnetic-bubble memory device includes the following steps: (1) formation of a conductor film via a 1st spacer (e.g., Al₂O₃) on a magnetic medium; (2) selective removal of the conductor film to form regions for a pattern of soft magnetic film for detection; (3) successive deposition of a soft magnetic film (e.g., Permalloy) and an insulator film (e.g., SiO₂) without removing the mask used in selective removal of the conductor film; (4) selective removal of the soft magnetic and insulator films by left off; (5) formation of the detection and conductor pattern by etching; (6) formation of a 2nd organic-insulator spacer (e.g., polyimide) over the extreme surfaces; (7) and formation of a soft-magnetic-film pattern for transferring. A device with planar surfaces and a 2-layer passivation insulator film is obtained.

IT Polyimides, uses and miscellaneous
(spacers, for bubble-domain memory devices)

L148 ANSWER 4 OF 4 HCPLUS COPYRIGHT ACS on STN

AN 1972:494621 HCPLUS Full-text

DN 77:94621

ED Entered STN: 12 May 1984

TI Semiconductor device having a passivation film and insulating films on a semiconductor substrate

PA Hitachi, Ltd.

IC H01L

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	GB 1283769	A	19720802	GB 1969-49538	19691008
PRAI	JP 1968-73134	A	19681009		

AB The amount of **surface** charge induced on a substrate **surface** by an elec. **passivation** film is controlled to a desired value by the insulating films produced on the **passivation** film. E.g., a SiO₂ film is formed on a p-type Si **substrate** and a SiO₂/P2O₅ film formed over the SiO₂ film. A SiO₂/B2O₃ film is then deposited over the SiO₂/P2O₅ film. Electrodes are provided on the **insulating** film and on the **substrate** to form a metal- **oxide**-semiconductor element. The **surface** charge d. on the **substrate** just under the SiO₂ film is $5 + 10^{11}/\text{cm}^2$. When a SiO₂/P2O₅ film is deposited on the SiO₂/B2O₃ film, the amount of **surface** charge induced on the Si **surface** is $2 + 10^{11}/\text{cm}^2$. A quadruple or quintuple layer has an even lower **surface** charge d. than the **triple** layer.

IT Semiconductor devices
(coating of, with multilayer insulating films)

IT Electric insulators and Dielectrics
(multilayer coatings, on semiconductor devices)

IT Coating materials
(multilayer, on semiconductor devices)

26/9/9

DIALOG(R) File 2:INSPEC

(c) Institution of Electrical Engineers. All rts. reserv.

00119652 INSPEC Abstract Number: A70023377, B70014091

Title: Oxidized SiH₄ as a diffusion source

Author(s): Mecs, B.M.

Author Affiliation: Stevens Inst. Technol., Hoboken, NJ, USA

Conference Title: 133rd meeting of the Electrochemical Society, Abstracts

p.15

Publisher: Electrochem. Soc, New York, NY, USA

Publication Date: 1968 Country of Publication: USA 158 pp.

Conference Sponsor: Electrochem. Soc

Conference Date: 5-9 May 1968 Conference Location: Boston, MA, USA

Language: English Document Type: Conference Paper (PA)

Abstract: Abstract only given. The mixture of silane (SiH₄) and phosphine (PH₃) in the presence of oxygen decomposes, and forms at or above 200 degrees C a continuous adherent layer of doped silicon dioxide on a silicon substrate. This oxide serves as a diffusion source. The apparatus required for deposition is simple. The effects of the deposition variables on the diffusion parameters can be independently determined. The results show that the useful range of surface concentration attainable extends from 5*10¹⁵ cm⁻³ to the solid solubility limit. The deposited film is also attractive as a passivation and/or insulation layer. Further by substituting diborane (B₂H₆) for phosphine a diffusion source for boron is obtainable.